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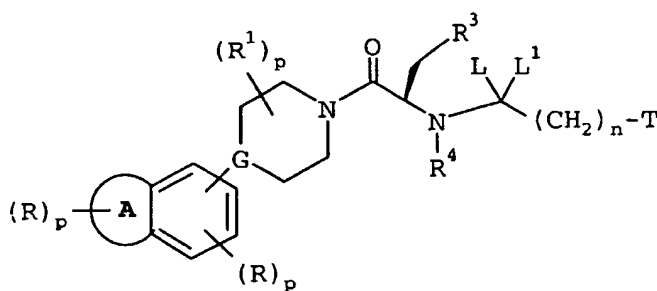
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(54) Title: SUBSTITUTED PIPERIDINES/PIPERAZINES AS MELANOCORTIN RECEPTOR AGONISTS



(I)

(57) Abstract: The present invention relates to melanocortin receptor agonists of formula I, which is useful in the treatment of obesity, diabetes and male and/or female sexual dysfunction.

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SUBSTITUTED PIPERIDINES/PIPERAZINES AS MELANOCORTIN RECEPTOR AGONISTS

FIELD OF THE INVENTION

The present invention relates to melanocortin receptor agonists, and more particularly piperazine and piperidine derivatives as melanocortin receptor agonists, which are useful for the treatment or prevention of diseases and disorders responsive to the activation of melanocortin receptors.

BACKGROUND OF THE INVENTION

Pro-opiomelanocortin (POMC) derived peptides are known to affect food intake. Several lines of evidence support the notion that the G-protein coupled receptors (GPCRs) of the melanocortin receptor (MC-R) family, several of which are expressed in the brain, are targets of POMC derived peptides involved in the control of food intake and metabolism.

Evidence for the involvement of MC-R in obesity includes: i) the agouti (A^{VY}) mouse which ectopically expresses an antagonist of the MC-1R, MC-3R and MC-4R is obese, indicating that blocking the action of these three MC-Rs can lead to hyperphagia and metabolic disorders; ii) MC-4R knockout mice (Huszar et al., *Cell*, 88:131-141, 1997) recapitulate the phenotype of the agouti mouse and these mice are obese; iii) the cyclic heptapeptide MC-1R, MC-3R, MC-4R, and MC-5R agonist melanotanin-II (MT-II) injected intracerebroventricularly (ICV) in rodents, reduces food intake in several animal feeding models (NPY, *ob/ob*, agouti, fasted) while ICV injected SHU-9119 (MC-3R, MC-4R antagonist;

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5 MC-1R and MC-5R agonist) reverses this effect and can induce hyperphagia; and iv) chronic intraperitoneal treatment of Zucker fatty rats with an alpha-NDP-MSH derivative (HP228) has been reported to activate MC-1R, MC-3R, MC-4R and MC-5R and to attenuate food intake and body weight gain over a 12 week period.

Five MC-Rs have thus far been identified, and these are expressed in
10 different tissues. MC-1R was initially characterized by dominant gain of function mutations at the extension locus, affecting coat color by controlling phaeomelanin to eumelanin conversion through control of tyrosinase. MC-1R is mainly expressed in melanocytes. MC-2R is expressed in the adrenal gland and represents the ACTH receptor. MC-3R is expressed in the brain, gut and placenta and may be involved in the
15 control of food intake and thermogenesis. MC-4R is uniquely expressed in the brain and its inactivation was shown to cause obesity. (A. Kask, et al., "Selective antagonist for the melanocortin-4-receptor (HS014) increases food intake in free-feeding rats, *Biochem. Biophys. Res. Commun.*, 245:90-93, 1998). MC-5R is expressed in many tissues including white fat, placenta and exocrine glands. A low level of expression is also
20 observed in the brain. MC-5R knock out mice reveal reduced sebaceous gland lipid production (Chen et al., *Cell*, 91:789-798, 1997).

MC-4R appears to play a role in other physiological functions as well, namely controlling grooming behavior, erection and blood pressure. Erectile dysfunction denotes the medical condition of inability to achieve penile erection sufficient for
25 successful intercourse. The term "impotence" is often times employed to describe this prevalent condition. Synthetic melanocortin receptor agonists have been found to initiate erections in men with psychogenic erectile dysfunction (H. Wessells et al., "Synthetic Melanotropic Peptide Initiates Erections in Men With Psychogenic Erectile Dysfunction: Double-Blind, Placebo Controlled Crossover Study," *J. Urol.*, 160: 389-393, 1998).
30 Activation of melanocortin receptors of the brain appears to cause normal stimulation of sexual arousal. Evidence for the involvement of MC-R in male and/or female sexual dysfunction is detailed in WO 00/74679.

Diabetes is a disease in which a mammal's ability to regulate glucose levels in the blood is impaired because the mammal has a reduced ability to convert
35 glucose to glycogen for storage in muscle and liver cells. In Type I diabetes, this reduced

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5 ability to store glucose is caused by reduced insulin production. "Type II Diabetes" or
"non-insulin dependent diabetes mellitus" (NIDDM) is the form of diabetes, which is due
to a profound resistance to insulin stimulating or regulatory effect on glucose and lipid
metabolism in the main insulin-sensitive tissues, muscle, liver and adipose tissue. This
resistance to insulin responsiveness results in insufficient insulin activation of glucose
10 uptake, oxidation and storage in muscle and inadequate insulin repression of lipolysis in
adipose tissue and of glucose production and secretion in liver. When these cells become
desensitized to insulin, the body tries to compensate by producing abnormally high levels
of insulin and hyperinsulemia results. Hyperinsulemia is associated with hypertension
and elevated body weight. Since insulin is involved in promoting the cellular uptake of
15 glucose, amino acids and triglycerides from the blood by insulin sensitive cells, insulin
insensitivity can result in elevated levels of triglycerides and LDL which are risk factors
in cardiovascular diseases. The constellation of symptoms which includes
hyperinsulemia combined with hypertension, elevated body weight, elevated triglycerides
and elevated LDL is known as Syndrome X.

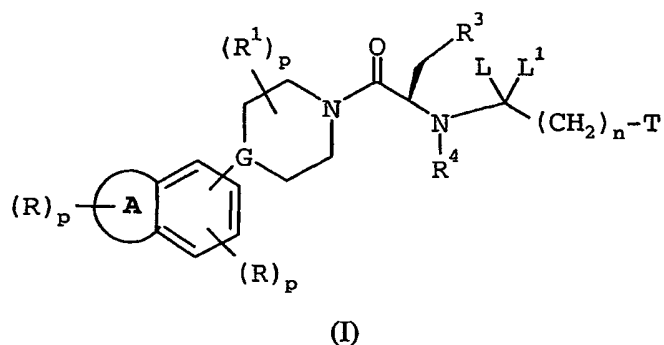
20 Spiropiperidine and piperidine derivates have been disclosed in U.S. Pat.
No. 6,294,534 B1, WO 01/70337, WO 00/74679 and WO 01/70708 as agonists of
melanocortin receptor(s), which can be used for the treatment of diseases and disorders,
such as obesity, diabetes and sexual dysfunction.

In view of the unresolved deficiencies in treatment of various diseases and
25 disorders as discussed above, it is an object of the present invention to provide novel
piperazine and piperidine derivatives, which are useful as melanocortin receptor agonists
to treat obesity, diabetes, and male and female sexual dysfunction.

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5 SUMMARY OF THE INVENTION

The present invention relates to a compound of novel piperazine and piperidine derivatives as melanocortin receptor agonists as shown formula I,



or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

G is CR¹ or N;

15 L and L¹ are independently hydrogen or together oxo;

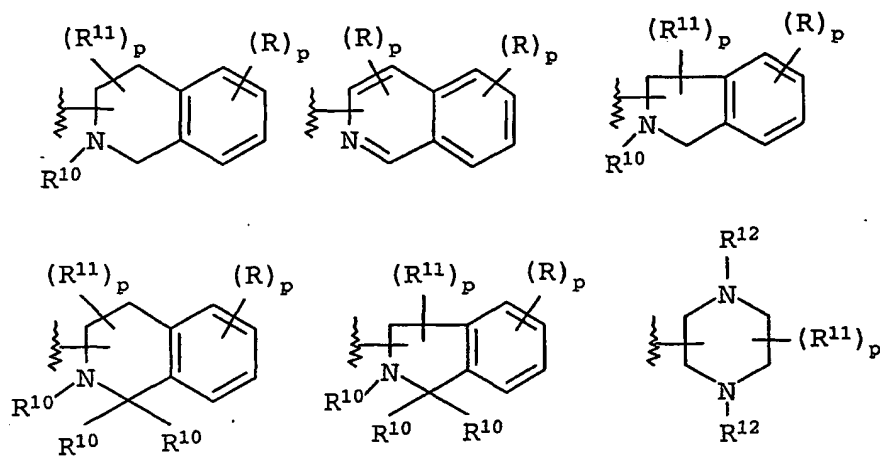


is a 5- or 6-membered carbocyclyl wherein the carbocyclyl being saturated, partially saturated or aromatic ring, and the carbocyclyl being optionally substituted with one to three substituents independently selected from R;

20

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5 T is:



R is independently:

- hydrogen,
- 10 hydroxy,
- (D)cyano,
- halo,
- C₁-C₈ alkyl,
- C₁-C₈ alkoxy,
- 15 C₃-C₇ cycloalkyl,
- C₁-C₄ haloalkyl,
- (D)heterocyclyl
- (D)C(O)R⁸,
- (D)C(O)(CH₂)_nN(R⁸)₂,
- 20 C₁-C₈ alkyl-N(R⁸)₂,
- (D)OR⁸,
- (D)OCOR⁸,
- (D)OC(O)N(R⁸)₂,
- (D)N(R⁸)₂,

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- 5 (D)NR⁸C(O)R⁸,
(D)NR⁸C(O)OR⁸,
(D)NR⁸C(O)N(R⁸)₂,
(D)NR⁸SO₂R⁸,
(D)SR⁸,
10 (D)SOR⁸,
(D)SO₂R⁸, or
(D)SO₂N(R⁸)₂;

R¹ is independently:

- 15 hydrogen, CONH(C₁-C₈ alkyl), C₁-C₈ alkyl, (D)phenyl, (D)C₃-C₇ cycloalkyl or
oxo, provided that oxo is not attached to the same carbon that attached to nitrogen
which forms an amide bond when G is N;

R³ is independently: aryl or thienyl;

- 20 wherein aryl and thienyl are optionally substituted with one to three substituents selected
from the group consisting of:
cyano, halo, C₁-C₈ alkyl, (D)C₃-C₇ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkyl
and C₁-C₄ haloalkyloxy;

- 25 R⁴ is independently:

hydrogen, C₁-C₈ alkyl, C(O)R⁸, C(O)OR⁸, C₃-C₇ cycloalkyl or (CH₂)_nO(C₁-C₈
alkyl), wherein n is 2-8;

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5 each R⁸ is independently:

hydrogen,

aryl, wherein aryl being phenyl or naphthyl,

C₁-C₈ alkyl,

C₁-C₄ alkoxy, or

10 (CH₂)_nC₁-C₄ haloalkyl, wherein n is 2-8;

each R¹⁰ is independently:

hydrogen, (C₁-C₈)alkyl, C(O)C₁-C₈ alkyl, aryl, or C₃-C₇ cycloalkyl;

15 each R¹¹ is independently:

hydrogen,

C₁-C₈ alkyl,

aryl,

heteroaryl

20 (CH₂)_nN(R⁸)₂,

(CH₂)_nNR⁸C(O)C₁-C₄ alkyl,

(CH₂)_nNR⁸SO₂C₁-C₄ alkyl,

(CH₂)_nSO₂N(R⁸)₂,

(CH₂)_n[O]_qC₁-C₈ alkyl,

25 (CH₂)_n[O]_q(CH₂)_nNR⁸COR⁸,

(CH₂)_n[O]_q(CH₂)_nNR⁸SO₂R⁸,

(CH₂)_n[O]_q-heterocyclyl or

(CH₂)_n[O]_q(C₁-C₈ alkyl)-heterocyclyl; and

wherein n is 2-8;

30

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- 5 each R¹² is independently:
hydrogen,
C₁-C₈ alkyl,
(D)C₃-C₇ cycloalkyl,
(D)phenyl
10 C(O)C₁-C₈ alkyl,
C(O)phenyl,
SO₂C₁-C₈ alkyl or
SO₂-phenyl;

- 15 D is a bond or -(CH₂)_n;
n is 0-8;
p is 0-4; and
q is 0-1.

20 The compounds of the present invention are useful in preventing or treating obesity or diabetes mellitus in a mammal comprising the administration of a therapeutically effective amount of the compound of formula I.

The compounds of the present invention are also useful in preventing or treating male or female sexual dysfunction in mammal, more specifically erectile dysfunction, comprising the administration of a therapeutically effective amount of the
25 compound of formula I.

Also within the scope of the present invention is a pharmaceutical composition or formulation which comprises a pharmaceutical carrier and at least one compound of formula I or its pharmaceutically acceptable salts or stereoisomers thereof.

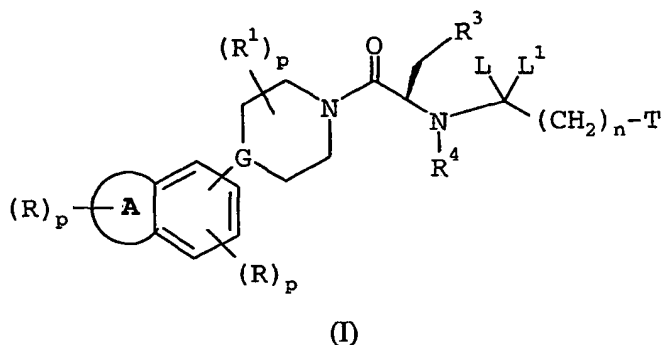
30 The present invention further includes a process of making a pharmaceutical composition or formulation comprising a compound of formula I or its pharmaceutically acceptable salt or stereoisomers thereof and a pharmaceutically acceptable carrier.

The present invention further includes a process of preparing a compound of formula I.

5 DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to melanocortin receptor agonists, and more particularly piperazine and piperidine derivatives as melanocortin receptor agonists. The compounds of present invention are useful for the treatment or prevention of diseases and disorders responsive to the activation of melanocortin receptors, such as obesity, diabetes and sexual dysfunction including erectile dysfunction and female sexual dysfunction.

An embodiment of the present invention is a compound of formula I,



or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

15

G is CR^1 or N;

L and L^1 are independently hydrogen or together oxo;

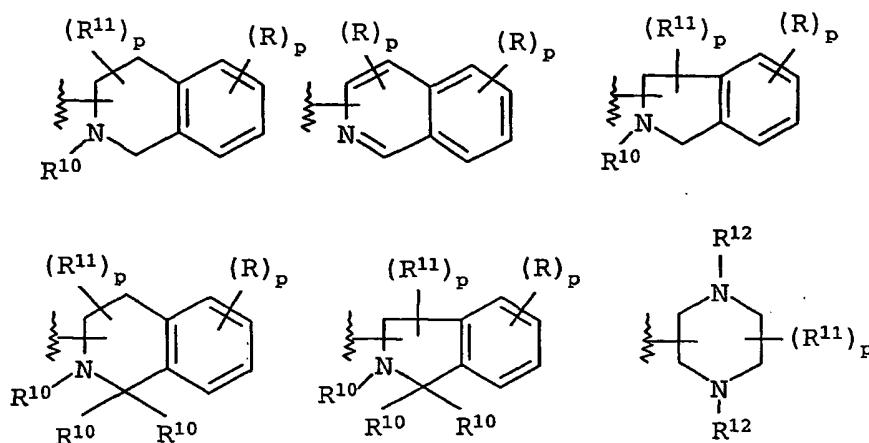
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is a 5- or 6-membered carbocyclyl wherein the carbocyclyl being saturated, partially saturated or aromatic ring, and the carbocyclyl being optionally substituted with one to three substituents independently selected from R;

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5 T is:



R is independently:

- hydrogen,
- hydroxy,
- 10 (D)cyano,
- halo,
- C_1 - C_8 alkyl,
- C_1 - C_8 alkoxy,
- C_3 - C_7 cycloalkyl,
- 15 C_1 - C_4 haloalkyl,
- (D)heterocyclyl
- (D) $C(O)R^8$,
- (D) $C(O)(CH_2)_nN(R^8)_2$,
- C_1 - C_8 alkyl- $N(R^8)_2$,
- 20 (D) OR^8 ,
- (D) $OCOR^8$,
- (D) $OC(O)N(R^8)_2$,
- (D) $N(R^8)_2$,
- (D) $NR^8C(O)R^8$,

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- 5 (D)NR⁸C(O)OR⁸,
 (D)NR⁸C(O)N(R⁸)₂,
 (D)NR⁸SO₂R⁸,
 (D)SR⁸,
 (D)SOR⁸,
 10 (D)SO₂R⁸, or
 (D)SO₂N(R⁸)₂;

R¹ is independently:

- hydrogen, CONH(C₁-C₈ alkyl), C₁-C₈ alkyl, (D)phenyl, (D)C₃-C₇ cycloalkyl or
 15 oxo, provided that oxo is not attached to the same carbon that attached to nitrogen
 which forms an amide bond when G is N;

R³ is independently: aryl or thienyl;

- wherein aryl and thienyl are optionally substituted with one to three substituents selected
 20 from the group consisting of:
 cyano, halo, C₁-C₈ alkyl, (D)C₃-C₇ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkyl
 and C₁-C₄ haloalkyloxy;

R⁴ is independently:

- 25 hydrogen, C₁-C₈ alkyl, C(O)R⁸, C(O)OR⁸, C₃-C₇ cycloalkyl or (CH₂)_nO(C₁-C₈
 alkyl), wherein n is 2-8;

each R⁸ is independently:

- hydrogen,
 30 aryl, wherein aryl being phenyl or naphthyl,
 C₁-C₈ alkyl,
 C₁-C₄ alkoxy, or

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5 $(\text{CH}_2)_n\text{C}_1\text{-C}_4$ haloalkyl, wherein n is 2-8;

each R^{10} is independently:

hydrogen, $(\text{C}_1\text{-C}_8)$ alkyl, $\text{C}(\text{O})\text{C}_1\text{-C}_8$ alkyl, aryl, or $\text{C}_3\text{-C}_7$ cycloalkyl;

10 each R^{11} is independently:

hydrogen,

$\text{C}_1\text{-C}_8$ alkyl,

aryl,

heteroaryl

15 $(\text{CH}_2)_n\text{N}(\text{R}^8)_2$,

$(\text{CH}_2)_n\text{NR}^8\text{C}(\text{O})\text{C}_1\text{-C}_4$ alkyl,

$(\text{CH}_2)_n\text{NR}^8\text{SO}_2\text{C}_1\text{-C}_4$ alkyl,

$(\text{CH}_2)_n\text{SO}_2\text{N}(\text{R}^8)_2$,

$(\text{CH}_2)_n[\text{O}]_q\text{C}_1\text{-C}_8$ alkyl,

20 $(\text{CH}_2)_n[\text{O}]_q(\text{CH}_2)_n\text{NR}^8\text{COR}^8$,

$(\text{CH}_2)_n[\text{O}]_q(\text{CH}_2)_n\text{NR}^8\text{SO}_2\text{R}^8$,

$(\text{CH}_2)_n[\text{O}]_q$ -heterocyclyl or

$(\text{CH}_2)_n[\text{O}]_q(\text{C}_1\text{-C}_8 \text{ alkyl})$ -heterocyclyl; and

wherein n is 2-8;

25

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5 each R¹² is independently:

hydrogen,

C₁-C₈ alkyl,

(D)C₃-C₇ cycloalkyl,

(D)phenyl

10 C(O)C₁-C₈ alkyl,

C(O)phenyl,

SO₂C₁-C₈ alkyl or

SO₂-phenyl;

15 D is a bond or -(CH₂)_n;

n is 0-8;

p is 0-4; and

q is 0-1.

20 The compound of the present invention as recited above, wherein the 5- or 6- membered carbocyclyl is cyclohexyl, cyclopentyl, cyclohexenyl, cyclopentenyl or phenyl.

The compound of the present invention as recited above, wherein R is independently at each occurrence: hydrogen, halo, hydroxy, C₁-C₈ alkyl, C₁-C₈ alkoxy, C₃-C₇ cycloalkyl, (D)heterocyclyl, (D)C(O)R⁸, (D)OC(O)R⁸, (D)C(O)aryl, 25 (D)C(O)(CH₂)_nN(R⁸)₂, (D)OC(O)(CH₂)_nN(R⁸)₂, (D)C(O)C₁-C₄ alkoxy, (D)N(R⁸)₂, (D)NR⁸COR⁸, (D)NR⁸C(O)C₁-C₄ alkoxy, (D)SR⁸, (D)SO₂R⁸, (D)SO₂N(R⁸)₂ or (D)NR⁸SO₂R⁸ where R⁸ independently at each occurrence being hydrogen, C₁-C₈ alkyl, C₁-C₈ alkoxy or phenyl.

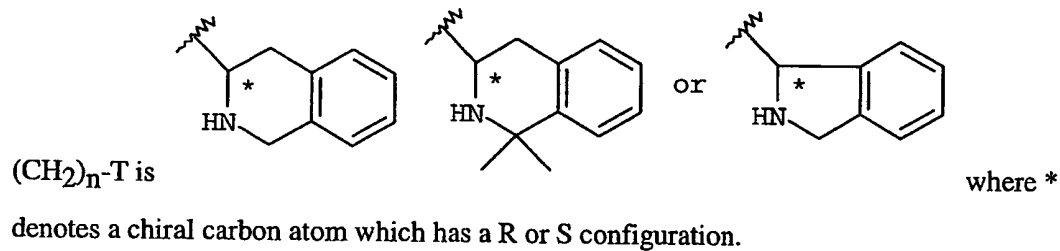
30 The compound of present invention as recited above, wherein R³ is phenyl optionally para-substituted with chloro, bromo, fluoro, methoxy, benzyloxy or methyl.

The preferred R³ is phenyl para-substituted with chloro, fluoro or methoxy.

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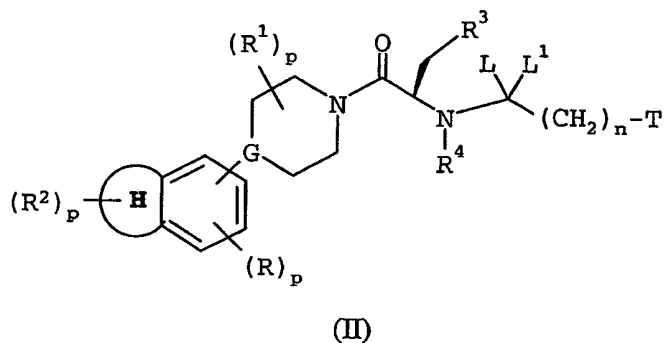
- 5 The compound of the present invention as recited above, wherein R^4 is hydrogen.

The compound of the present invention as recited above, wherein -



- 10 The compound of the present invention as recited above, wherein L and L¹ are together oxo and the chiral carbon has R configuration.

Another embodiment of the present invention is a compound of formula II,




- 15 or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

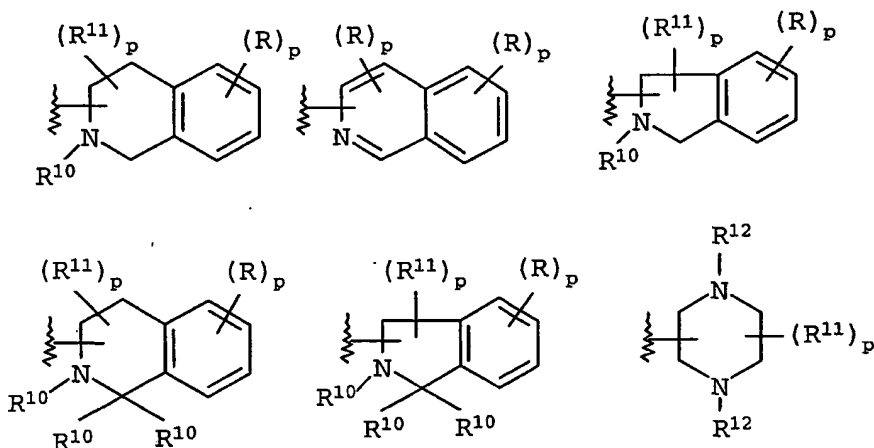
G is CR¹ or N;

L and L¹ are independently hydrogen or together oxo;

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- 5  is a 5- or 6-membered heterocyclyl,
 wherein the heterocyclyl being saturated, partially saturated or aromatic ring, provided that
 when G is carbon, the 5-membered nitrogen containing heterocyclyl does not contain a
 carbon-carbon double bond in the ring; and
 wherein the heterocyclyl being optionally substituted with one to three substituents
 10 independently selected from R²;

T is:



R is independently:

- 15 hydrogen,
 hydroxy,
 (D)cyano,
 halo,
 C₁-C₈ alkyl,
 20 C₁-C₈ alkoxy,
 C₃-C₇ cycloalkyl,
 C₁-C₄ haloalkyl,
 (D)heterocyclyl,
 (D)C(O)R⁸,

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- 5 (D)C(O)(CH₂)_nN(R⁸)₂,
 C₁-C₈ alkyl-N(R⁸)₂,
 (D)OR⁸,
 (D)OCOR⁸,
 (D)OC(O)N(R⁸)₂,
10 (D)N(R⁸)₂,
 (D)NR⁸C(O)R⁸,
 (D)NR⁸C(O)OR⁸,
 (D)NR⁸C(O)N(R⁸)₂,
 (D)NR⁸SO₂R⁸,
15 (D)SR⁸,
 (D)SOR⁸,
 (D)SO₂R⁸, or
 (D)SO₂N(R⁸)₂;
20 R¹ is independently:
 hydrogen, CONH(C₁-C₈ alkyl), C₁-C₈ alkyl, (D)phenyl, (D)C₃-C₇ cycloalkyl or
 oxo, provided that oxo is not attached to the same carbon that attached to nitrogen
 which forms an amide bond when G is N;

25

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- 5 R^2 is independently:
hydrogen,
C₁-C₈ alkyl,
C₃-C₇ cycloalkyl,
C(O)NH₂,
10 C(O)O(C₁-C₆ alkyl),
C(O)(CH₂)_nN(C₁-C₆ alkyl)₂,
C(O)-phenyl,
SO₂(C₁-C₆ alkyl),
SO₂N(C₁-C₆ alkyl)₂,
15 SO₂-phenyl or
C(O)(C₁-C₆ alkyl);

- R^3 is independently: aryl or thienyl;
wherein aryl and thienyl are optionally substituted with one to three substituents selected
20 from the group consisting of:
cyano, halo, C₁-C₈ alkyl, (D)C₃-C₇ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkyl
and C₁-C₄ haloalkyloxy;

- R^4 is independently:
25 hydrogen, C₁-C₈ alkyl, C(O)R⁸, C(O)OR⁸, C₃-C₇ cycloalkyl or (CH₂)_nO(C₁-C₈
alkyl), wherein n is 2-8;

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5 each R⁸ is independently:

hydrogen,

aryl, wherein aryl being phenyl or naphthyl,

C₁-C₈ alkyl,

C₁-C₄ alkoxy, or

10 (CH₂)_nC₁-C₄ haloalkyl, wherein n is 2-8;

each R¹⁰ is independently:

hydrogen, (C₁-C₈)alkyl, C(O)C₁-C₈ alkyl, aryl, or C₃-C₇ cycloalkyl;

15 each R¹¹ is independently:

hydrogen,

C₁-C₈ alkyl,

aryl,

heteroaryl

20 (CH₂)_nN(R⁸)₂,

(CH₂)_nNR⁸C(O)C₁-C₄ alkyl,

(CH₂)_nNR⁸SO₂C₁-C₄ alkyl,

(CH₂)_nSO₂N(R⁸)₂,

(CH₂)_n[O]_qC₁-C₈ alkyl,

25 (CH₂)_n[O]_q(CH₂)_nNR⁸COR⁸,

(CH₂)_n[O]_q(CH₂)_nNR⁸SO₂R⁸,

(CH₂)_n[O]_q-heterocyclyl or

(CH₂)_n[O]_q(C₁-C₈ alkyl)-heterocyclyl; and

wherein n is 2-8;

30

each R¹² is independently:

hydrogen,

-19-

- 5 C_1 - C_8 alkyl,
 (D) C_3 - C_7 cycloalkyl,
 (D)phenyl
 C(O) C_1 - C_8 alkyl,
 C(O)phenyl,
 10 SO_2C_1 - C_8 alkyl or
 SO_2 -phenyl;

D is a bond or $-(CH_2)_n$;

n is 0-8;

- 15 p is 0-4; and

q is 0-1.

The compound of present invention as recited above, wherein the 5- or 6-membered heterocyclc contains at least one nitrogen.

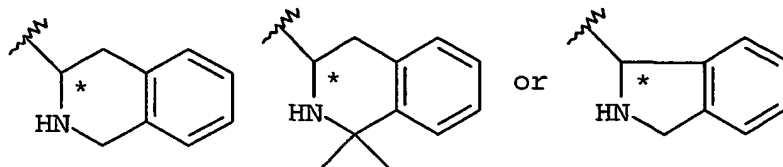
- 20 The compound of present invention as recited above, wherein the heterocyclcyl is piperidinyl, pyrrolidinyl, pyrrolinyl, isoxazolyl, oxazolyl, thiazoyl, triazolyl, tetrazolyl, thiadiazolyl or oxadiazolyl.

The compound of present invention as recited above, wherein R^3 is phenyl optionally para-substituted with chloro, bromo, fluoro, methoxy, benzyloxy or methyl.

- 25 The compound of present invention as recited above, wherein R^3 is phenyl para-substituted with chloro, fluoro or methoxy.

The compound of present invention as recited above, wherein R^4 is hydrogen.

The compound of present invention as recited above, wherein $-(CH_2)_n$ -T



is

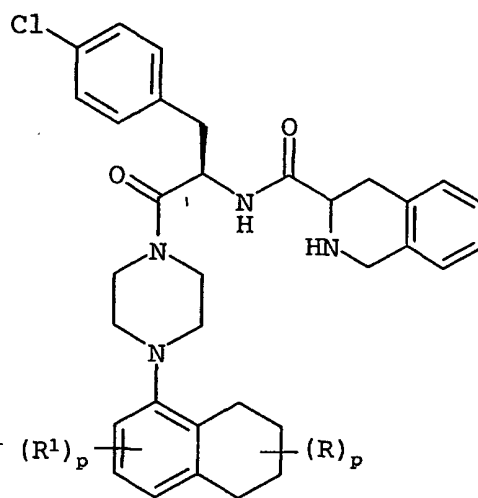
where * denotes a

- 30 chiral carbon atom which has a R or S configuration.

-20-

5 The compound of present invention as recited above, wherein L and L¹ are together oxo and the chiral carbon has R configuration.

 The preferred embodiment of the present invention provides a compound of formula III,



(III)

10 or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

R is selected from the group consisting of:

15 C₁-C₆ alkoxy, NH₂, NH(C₁-C₆ alkyl), N(C₁-C₆ alkyl)₂, NHC(O)O(C₁-C₆ alkyl), N(CH₃)SO₂(C₁-C₆ alkyl), NHSO₂(C₁-C₆ alkyl), NHC(O)(C₁-C₆ alkyl), OC(O)(C₁-C₆ alkyl), OC(O)NH₂, heterocyclyl, C₁-C₆ alkoxy, hydrogen and C₃-C₇ cycloalkyl;

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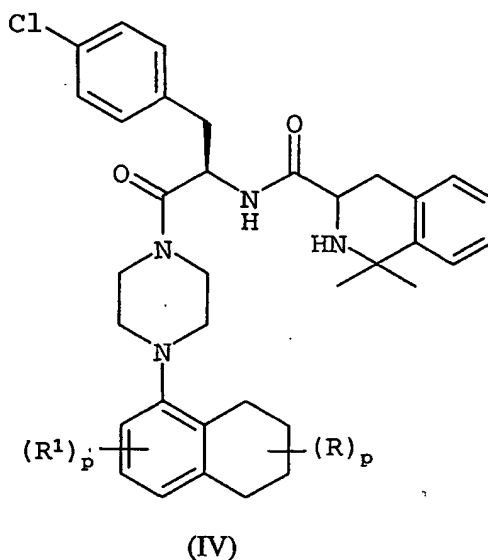
5 R^1 is selected from the group consisting of:

halo, C_1 - C_6 alkyl, $N(C_1$ - C_6 alkyl) $_2$, C_1 - C_4 haloalkyl, C_1 - C_6 alkoxy and hydrogen; and

p is 0-4.

Another preferred embodiment of the present invention provides a

10 compound of formula IV,



or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

15 R is selected from the group consisting of:

C_1 - C_6 alkoxy, NH_2 , $NH(C_1$ - C_6 alkyl), $N(C_1$ - C_6 alkyl) $_2$, $NHC(O)O(C_1$ - C_6 alkyl), $N(CH_3)SO_2(C_1$ - C_6 alkyl), $NHSO_2(C_1$ - C_6 alkyl), $NHC(O)(C_1$ - C_6 alkyl), $OC(O)(C_1$ - C_6 alkyl), $OC(O)NH_2$, heterocyclyl, C_1 - C_6 alkoxy, hydrogen and C_3 - C_7 cycloalkyl;

20 R^1 is selected from the group consisting of:

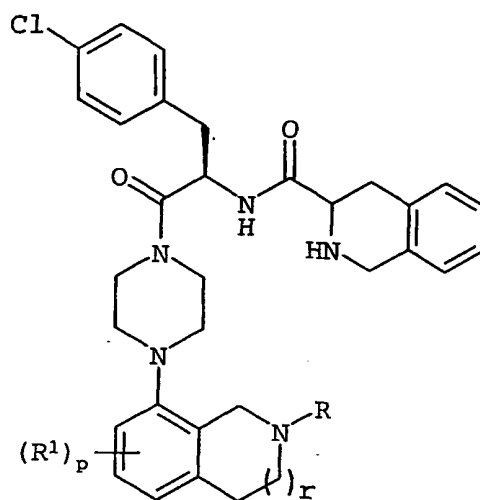
halo, C_1 - C_6 alkyl, $N(C_1$ - C_6 alkyl) $_2$, C_1 - C_4 haloalkyl, C_1 - C_6 alkoxy and hydrogen; and

p is 0-4.

Yet another preferred embodiment of the present invention provides a

25 compound of formula V,

-22-



(V)

or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

R is selected from the group consisting of:

- hydrogen, C₁-C₆ alkyl, C(O)NH₂, C(O)O(C₁-C₆ alkyl), C(O)(CH₂)_nN(C₁-C₆ alkyl)₂, C(O)-phenyl, SO₂(C₁-C₆ alkyl), SO₂N(C₁-C₆ alkyl)₂, SO₂-phenyl, C(O)C₁-C₆ alkyl, C₃-C₇ cycloalkyl and (C₂-C₆ alkyl)N(C₁-C₆ alkyl)₂;

R¹ is selected from the group consisting of:

halo, C₁-C₆ alkyl, N(C₁-C₆ alkyl)₂, C₁-C₄ haloalkyl, hydrogen and C₁-C₆ alkoxy;

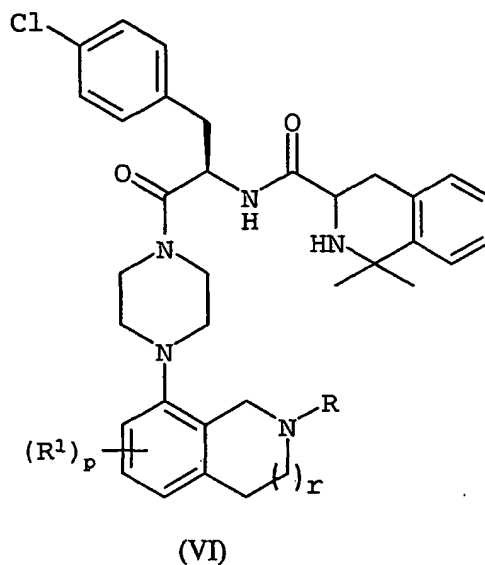
n is 0-8

p is 0-4; and

r is 0-1.

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- 5 Yet another preferred embodiment of the present invention provides a compound of formula VI,



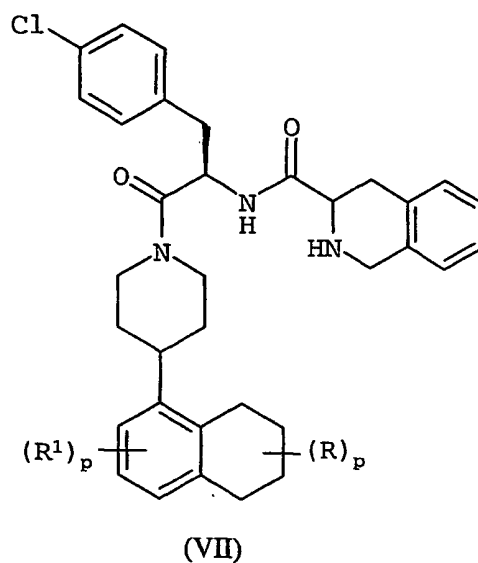
or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

- 10 R is selected from the group consisting of:
hydrogen, C₁-C₆ alkyl, C(O)NH₂, C(O)O(C₁-C₆ alkyl), C(O)(CH₂)_nN(C₁-C₆
alkyl)₂, C(O)-phenyl, SO₂(C₁-C₆ alkyl), SO₂N(C₁-C₆ alkyl)₂, SO₂-phenyl,
C(O)C₁-C₆ alkyl, C₃-C₇ cycloalkyl and (C₂-C₆ alkyl)N(C₁-C₆ alkyl)₂;
R¹ is selected from the group consisting of:
15 halo, C₁-C₆ alkyl, N(C₁-C₆ alkyl)₂, C₁-C₄ haloalkyl, hydrogen and C₁-C₆
alkoxy;
n is 0-8
p is 0-4; and
r is 0-1.

20

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- 5 Yet another preferred embodiment of the present invention provides a compound of formula VII,

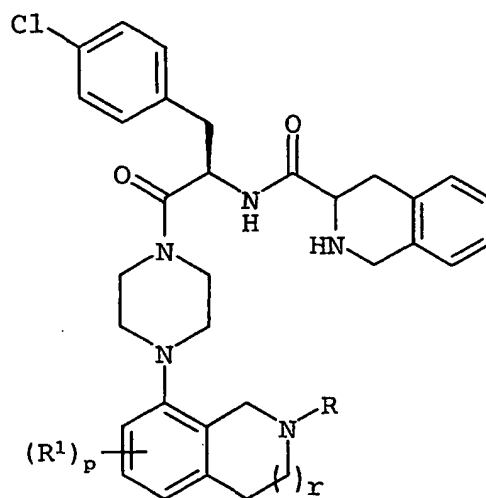


or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

- 10 R is selected from the group consisting of:
 C_1 - C_6 alkoxy, NH_2 , $NH(C_1-C_6 \text{ alkyl})$, $N(C_1-C_6 \text{ alkyl})_2$, $NHC(O)O(C_1-C_6 \text{ alkyl})$, $N(CH_3)SO_2(C_1-C_6 \text{ alkyl})$, $NHSO_2(C_1-C_6 \text{ alkyl})$, $NHC(O)(C_1-C_6 \text{ alkyl})$, $OC(O)(C_1-C_6 \text{ alkyl})$, $OC(O)NH_2$, heterocyclyl, C_1 - C_6 alkoxy, hydrogen and C_3 - C_7 cycloalkyl;
- 15 R^1 is selected from the group consisting of:
halo, C_1 - C_6 alkyl, $N(C_1-C_6 \text{ alkyl})_2$, C_1 - C_4 haloalkyl, hydrogen and C_1 - C_6 alkoxy; and
- p is 0-4.

-25-

5 Yet another preferred embodiment of the present invention provides a compound of formula VIII,



(VIII)

or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

10 R is selected from the group consisting of:

hydrogen, C₁-C₆ alkyl, C(O)NH₂; C(O)O(C₁-C₆ alkyl), C(O)(CH₂)_nN(C₁-C₆ alkyl)₂, C(O)-phenyl, SO₂(C₁-C₆ alkyl), SO₂N(C₁-C₆ alkyl)₂, SO₂-phenyl, C(O)C₁-C₆ alkyl, C₃-C₇ cycloalkyl and (C₂-C₆ alkyl)N(C₁-C₆ alkyl)₂;

R^1 is selected from the group consisting of:

15 halo, C₁-C₆ alkyl, N(C₁-C₆ alkyl)₂, C₁-C₄ haloalkyl, hydrogen and C₁-C₆ alkoxy;

n is 0-8

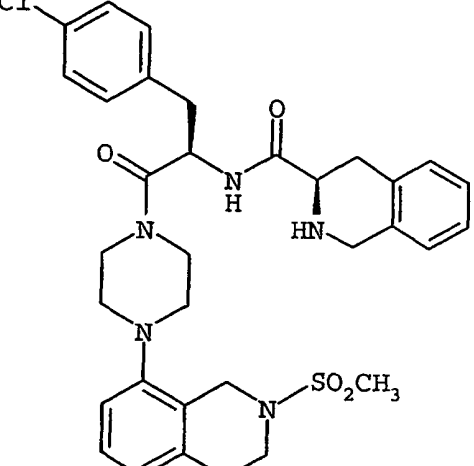
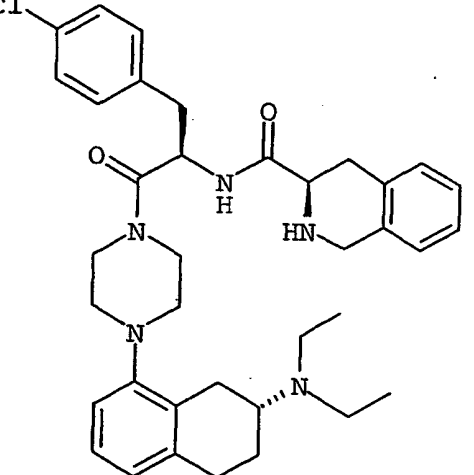
p is 0-4; and

¹ r is 0-1.

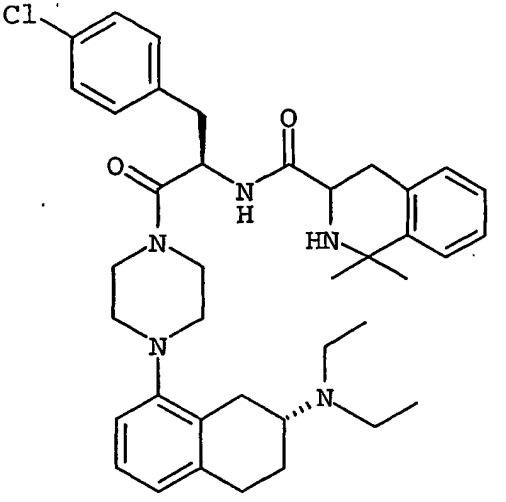
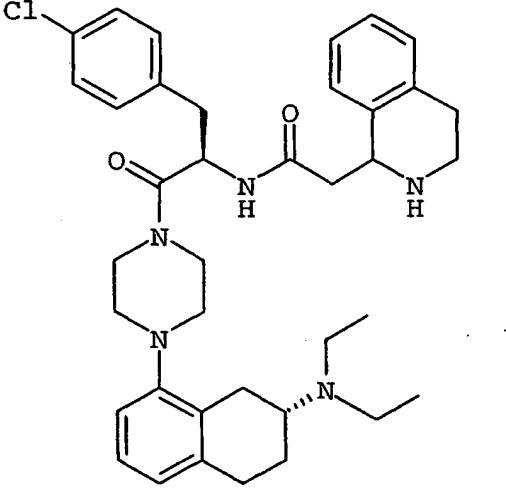
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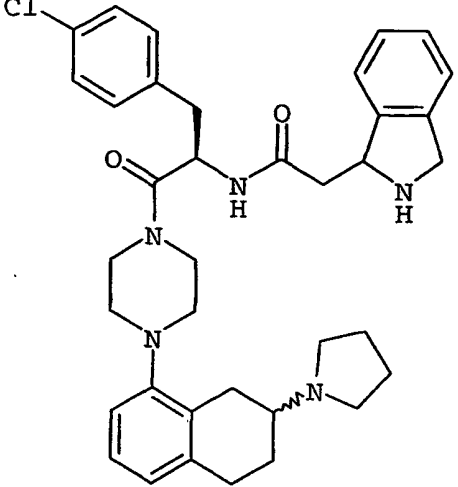
- 5 The most preferred compound of the present invention is selected from the group consisting of:

Name of compound	Structure of compound
1,2,3,4-tetrahydro-isoquinoline-3-carboxylic acid {1-(4-chlorobenzyl)-2-[4-(2-methanesulfonyl-1,2,3,4-tetrahydro-isoquinoline-8-yl)-piperazin-1-yl]-2-oxo-ethyl}-amide	
1,2,3,4-tetrahydro-isoquinoline-3-carboxylic acid {1-(4-chlorobenzyl)-2-[4-(7-diethylamino-5,6,7,8-tetrahydronaphthalen-1-yl)-piperazin-1-yl]-2-oxo-ethyl}-amide	

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Name of compound	Structure of compound
<p>1,1-dimethyl-1,2,3,4-tetrahydro-isoquinoline-3-carboxylic acid {1-(4-chlorobenzyl)-2-[4-(7-diethylamino-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-2-oxo-ethyl}-amide</p>	 <p>and its stereoisomers</p>
<p>N-{1-(4-chloro-benzyl)-2-[4-(7-diethylamino-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-2-oxo-ethyl}-2-(1,2,3,4-tetrahydro-isoquinolin-1-yl)-acetamide</p>	 <p>and its stereoisomers</p>

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Name of compound	Structure of compound
N-{1-(4-chloro-benzyl)-2-oxo-2-[4-(7-pyridin-1-yl-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-ethyl}-2-(2,3-dihydro-1H-isindol-1-yl)-acetamide	 <p>and its stereoisomers</p>

5

Also encompassed by the present invention is a pharmaceutical composition or formulation, which comprises a pharmaceutical carrier and at least one compound of formula I or its pharmaceutically acceptable salts or stereoisomers thereof. The pharmaceutical composition and or formulation may optionally further include a second active ingredient selected from the group consisting of an insulin sensitizer, insulin mimetic, sulfonylurea, alpha-glycosidase inhibitor, HMG-CoA reductase inhibitor, sequestrant cholesterol lowering agent, beta 3 adrenergic receptor agonist, neuropeptide Y antagonist, phosphodiester V inhibitor, and an alpha 2 adrenergic receptor antagonist.

10

Yet another aspect of the present invention is a process of making a pharmaceutical composition comprising a compound of formula I or its pharmaceutically acceptable salt or stereoisomers thereof as recited above and a pharmaceutically acceptable carrier.

Yet another aspect of the present invention is a method of preventing or treating obesity or diabetes mellitus in mammal comprising the administration of a therapeutically effective amount of the compound of formula I.

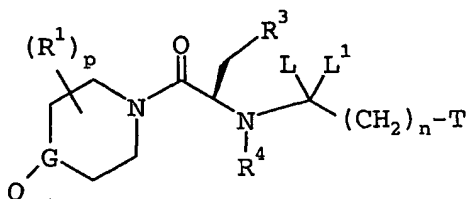
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Yet another aspect of the present invention is a method of preventing or treating male or female sexual dysfunction in mammal, more specifically the male or

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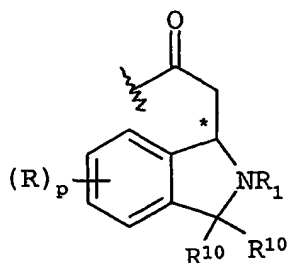
- 5 female sexual dysfunction, comprising the administration of a therapeutically effective amount of the compound of formula I.

Yet another aspect of the present invention is a process for preparing a compound of formula I:



- 10 or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

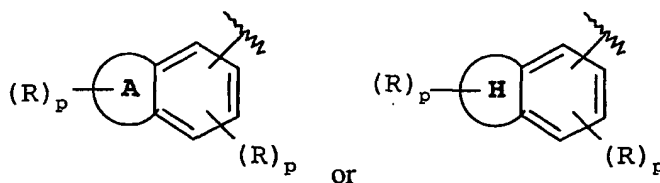
-CLL¹-(CH₂)ₙ-T is:



wherein R₁ is hydrogen, C₁-C₈ alkyl, Boc, CBZ, FMOC, phenyl or (C₁-C₈ alkyl)phenyl;

15

Q represents a moiety:




G is CR¹ or N;



- 20 is a 5- or 6-membered carbocyclyl wherein the carbocyclyl being saturated, partially saturated or aromatic ring, and the carbocyclyl being optionally substituted with one to three substituents independently selected from R;

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- 5  is a 5- or 6-membered heterocyclyl,
 wherein the heterocyclyl being saturated, partially saturated or aromatic ring, provided that
 when G is carbon, the 5-membered nitrogen containing heterocyclyl does not contain a
 carbon-carbon double bond in the ring; and
 wherein the heterocyclyl being optionally substituted with one to three substituents
 10 independently selected from R²;

R is independently:

- hydrogen,
 hydroxy,
 15 (D)cyano,
 halo,
 C₁-C₈ alkyl,
 C₁-C₈ alkoxy,
 C₃-C₇ cycloalkyl,
 20 C₁-C₄ haloalkyl,
 (D)heterocyclyl
 (D)C(O)R⁸,
 (D)C(O)(CH₂)_nN(R⁸)₂,
 C₁-C₈ alkyl-N(R⁸)₂,
 25 (D)OR⁸,
 (D)OCOR⁸,
 (D)OC(O)N(R⁸)₂,
 (D)N(R⁸)₂,
 (D)NR⁸C(O)R⁸,
 30 (D)NR⁸C(O)OR⁸,
 (D)NR⁸C(O)N(R⁸)₂,

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- 5 (D)NR⁸SO₂R⁸,
(D)SR⁸,
(D)SOR⁸,
(D)SO₂R⁸, or
(D)SO₂N(R⁸)₂;

10

R¹ is independently:

hydrogen, CONH(C₁-C₈ alkyl), C₁-C₈ alkyl, (D)phenyl, (D)C₃-C₇ cycloalkyl or
oxo, provided that oxo is not attached to the same carbon that attached to nitrogen
which forms an amide bond when G is N;

15

R² is independently:

hydrogen,
C₁-C₈ alkyl,
C₃-C₇ cycloalkyl,
20 C(O)NH₂,
C(O)O(C₁-C₆ alkyl),
C(O)(CH₂)_nN(C₁-C₆ alkyl)₂,
C(O)-phenyl,
SO₂(C₁-C₆ alkyl),
25 SO₂N(C₁-C₆ alkyl)₂,
SO₂-phenyl or
C(O)(C₁-C₆ alkyl);

R³ is independently: aryl or thienyl;

- 30 wherein aryl and thienyl are optionally substituted with one to three substituents selected
from the group consisting of:

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5 cyano, halo, C₁-C₈ alkyl, (D)C₃-C₇ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkyl
and C₁-C₄ haloalkyloxy;

R⁴ is independently:

hydrogen, C₁-C₈ alkyl, C(O)R⁸, C(O)OR⁸, C₃-C₇ cycloalkyl or (CH₂)_nO(C₁-C₈
10 alkyl), wherein n is 2-8;

each R⁸ is independently:

hydrogen,
aryl, wherein aryl being phenyl or naphthyl,
15 C₁-C₈ alkyl,
C₁-C₄ alkoxy, or
(CH₂)_nC₁-C₄ haloalkyl, wherein n is 2-8;

each R¹⁰ is independently:

20 hydrogen, (C₁-C₈)alkyl, C(O)C₁-C₈ alkyl, aryl, or C₃-C₇ cycloalkyl;

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5 D is a bond or $-(CH_2)_n-$;

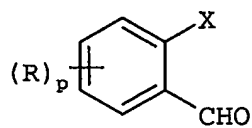
n is 0-8;

p is 0-4; and

q is 0-1;

10 comprising the steps of:

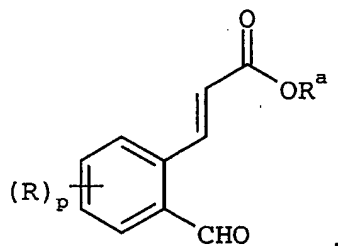
a) reacting a compound having a structural formula 1,



(1)

with $CH_2CH=C(O)OR^a$ wherein R^a is hydrogen or C_1 - C_8 alkyl and X is halo, in the

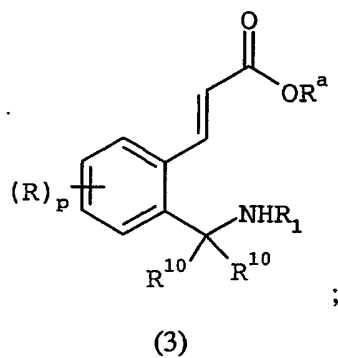
15 presence of a catalyst and a base in a suitable organic solvent to give the compound of formula 2,



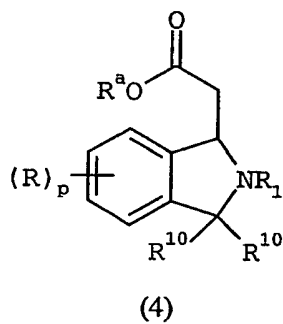
(2)

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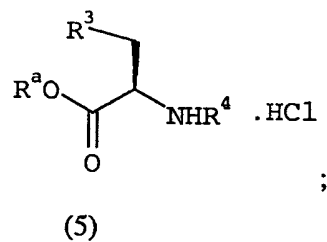
- 5 b) reductively aminating the compound of formula 2 in the presence of amine in an acidic condition to give a compound of formula 3,



- 10 c) cyclizing the compound of formula 3 by Michael addition to give a compound of formula 4 or stereoisomers thereof,

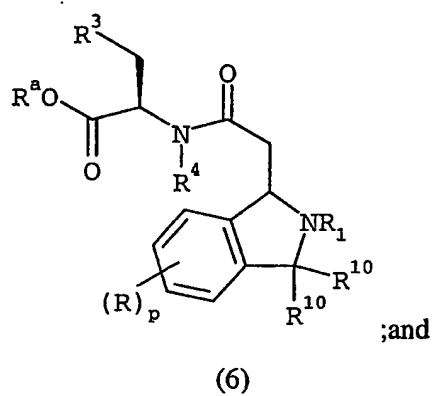


- 15 d) coupling the compound of formula 4 or stereoisomers thereof, wherein R^a of compound 4 is H, with a compound of formula 5,

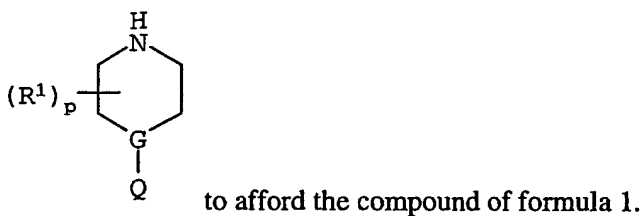


-35-

- 5 wherein R^a of compound 5 is C₁-C₈ alkyl, to give a compound of formula 6;



- 10 e) coupling the compound of formula 6, wherein R^a is H, with a compound having a structural,



The process of the present invention as recited above, wherein

- 15 in Step (a) is 2-bromobenzaldehydes.
- The chemical structure shows a benzene ring with a substituent (R)_p and a group X. The ring is also substituted with a CHO group.

The process of the present invention as recited above, wherein CH₂CH=C(O)OR in Step (a) is methylacrylate.

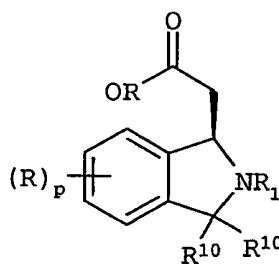
- The process of the present invention as recited above, wherein the catalyst in Step (a) is selected from the group consisting of: Pd(Ph₃P)₂Cl₂, Pd(Ph₃P)₄Cl₂,
 20 Pd(Ph₃P)₄, Pd(Ph₃P)₂Cl₂/CuI, Pd(OAc)₂/Ph₃P-Bu₄NBr, Pd(Ph₃P)₄Cl₂/H₂ and Pd(OAc)₂/P(O-tol)₃; and wherein the base in Step (a) is NR₃ wherein R is hydrogen or C₁-C₈ alkyl.

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5 The process of the present invention as recited above, wherein the amine in Step (b) is selected from the group consisting of: benzylamine, alpha-methylbenzylamine and BocNH₂.

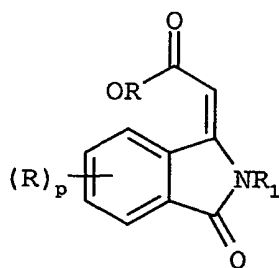
 The process of the present invention as recited above, wherein the Step (b) further comprises reducing of intermediate imine compound in the presence of reducing
10 agent, the reducing agent being selected from the group consisting of: NaCNBH₃, Na(OAc)₃BH, NaBH₄/H⁺, and a combination of Et₃SiH and TFA in CH₃CN or CH₂Cl₂.

 The process of the present invention as recited above, wherein the stereoisomer of compound of formula 4 in Step (c) is a compound of formula 4a.



(4a)

 The process of the present invention as recited above, wherein the compound of formula 4a is prepared by asymmetric hydrogenation of a compound having structural formula,



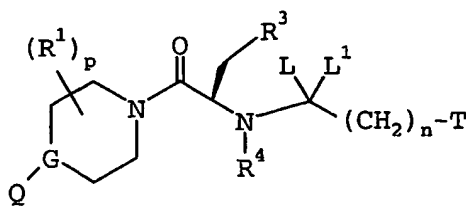
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 The process of the present invention as recited above, wherein the Michael addition in Step (c) is carried out in a basic workup condition.

 The process of the present invention as recited above, wherein the Step (e) further comprises deprotecting or protecting of the compound of formula (4) at NR₁.

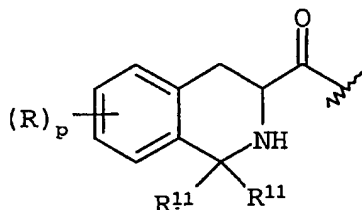
-37-

- 5 Yet another aspect of the present invention is a process for preparing a compound of formula I:

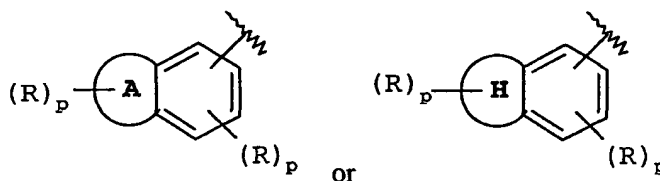


(I)

- 10 or a pharmaceutically acceptable salts or stereoisomers thereof, wherein -CLL¹-(CH₂)ₙ-T is:



Q represents a moiety:



- 15 G is CR¹ or N;



is a 5- or 6-membered carbocyclyl wherein the carbocyclyl being saturated, partially saturated or aromatic ring, and the carbocyclyl being optionally substituted with one to three substituents independently selected from R;

20



is a 5- or 6-membered heterocyclyl,

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- 5 wherein the heterocyclyl being saturated, partially saturated or aromatic ring, provided that when G is carbon, the 5-membered nitrogen containing heterocyclyl does not contain a carbon-carbon double bond in the ring; and wherein the heterocyclyl being optionally substituted with one to three substituents independently selected from R²;

10

R is independently:

- hydrogen,
hydroxy,
(D)cyano,
15 halo,
C₁-C₈ alkyl,
C₁-C₈ alkoxy,
C₃-C₇ cycloalkyl,
C₁-C₄ haloalkyl,
20 (D)heterocyclyl
(D)C(O)R⁸,
(D)C(O)(CH₂)_nN(R⁸)₂,
C₁-C₈ alkyl-N(R⁸)₂,
(D)OR⁸,
25 (D)OCOR⁸,
(D)OC(O)N(R⁸)₂,
(D)N(R⁸)₂,
(D)NR⁸C(O)R⁸,
(D)NR⁸C(O)OR⁸,
30 (D)NR⁸C(O)N(R⁸)₂,
(D)NR⁸SO₂R⁸,
(D)SR⁸,

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- 5 (D)SOR⁸,
(D)SO₂R⁸, or
(D)SO₂N(R⁸)₂;

R¹ is independently:

- 10 hydrogen, CONH(C₁-C₈ alkyl), C₁-C₈ alkyl, (D)phenyl, (D)C₃-C₇ cycloalkyl or
oxo, provided that oxo is not attached to the same carbon that attached to nitrogen
which forms an amide bond when G is N;

R² is independently:

- 15 hydrogen,
C₁-C₈ alkyl,
C₃-C₇ cycloalkyl,
C(O)NH₂,
C(O)O(C₁-C₆ alkyl),
20 C(O)(CH₂)_nN(C₁-C₆ alkyl)₂,
C(O)-phenyl,
SO₂(C₁-C₆ alkyl),
SO₂N(C₁-C₆ alkyl)₂,
SO₂-phenyl or
25 C(O)(C₁-C₆ alkyl);

-40-

- 5 R^3 is independently: aryl or thienyl;
wherein aryl and thienyl are optionally substituted with one to three substituents selected from the group consisting of:
 cyano, halo, C_1 - C_8 alkyl, (D) C_3 - C_7 cycloalkyl, C_1 - C_4 alkoxy, C_1 - C_4 haloalkyl
 and C_1 - C_4 haloalkyloxy;

10

-41-

- 5 R^4 is independently:
hydrogen, C_1 - C_8 alkyl, $C(O)R^8$, $C(O)OR^8$, C_3 - C_7 cycloalkyl or $(CH_2)_nO(C_1$ - C_8
alkyl), wherein n is 2-8;

each R^8 is independently:

- 10 hydrogen,
aryl, wherein aryl being phenyl or naphthyl,
 C_1 - C_8 alkyl,
 C_1 - C_4 alkoxy, or
 $(CH_2)_nC_1$ - C_4 haloalkyl, wherein n is 2-8;

- 15 each R^{11} is independently: hydrogen or $(C_1$ - $C_8)$ alkyl;

D is a bond or $-(CH_2)_n-$;

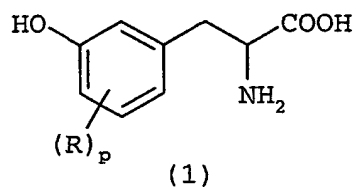
n is 0-8;

- 20 p is 0-4; and

q is 0-1;

comprising the steps of:

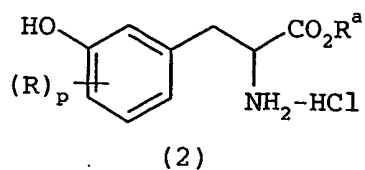
- a) esterifying a compound of formula 1,



25

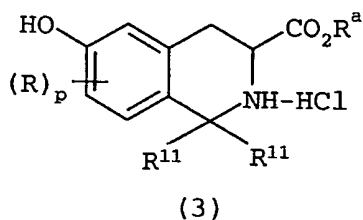
-42-

- 5 with an alcohol R^aOH to form a compound of formula 2,



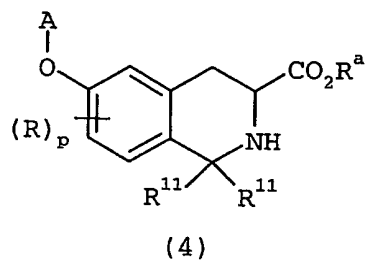
wherein R^a is C_1 - C_4 alkyl or (D)phenyl;

- 10 b) reacting a compound of formula 2 with $R^{11}COR^{11}$ to form a compound of formula 3,



wherein R^{11} is independently hydrogen or C_1 - C_4 alkyl;

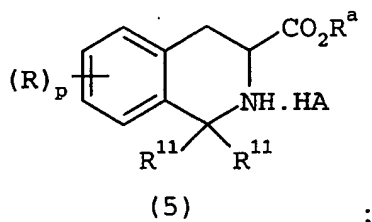
- 15 c) reacting a compound of formula 3 with an activating group to form a compound of formula 4,



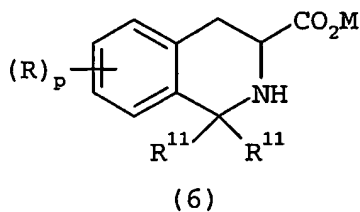
wherein A is an activating group;

-43-

- 5 d) deoxygenating the compound of formula 4 by hydrogenation to afford a compound of formula 5,

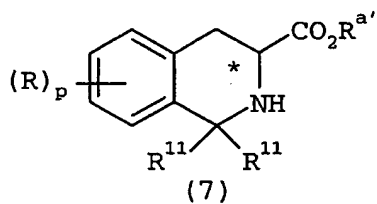


- 10 e) optionally reacting the compound of formula 5 with an inorganic base to form a compound of formula 6,



wherein HA is an acidic and M is a univalent cation;

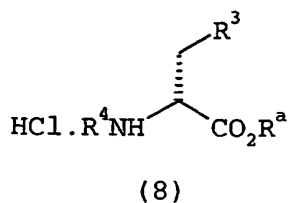
- 15 f) resolving the compound of formula 5 or formula 6 to afford a chiral compound of formula 7,



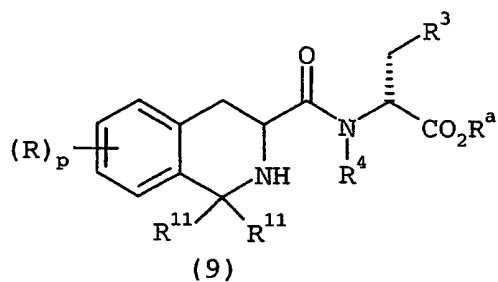
wherein M is hydrogen and R^a is H or R^a;

-44-

- 5 g) coupling the compound of formula 7 with a compound of formula 8,

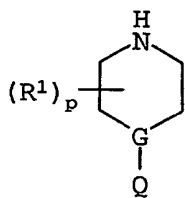


to afford a compound of formula 9,



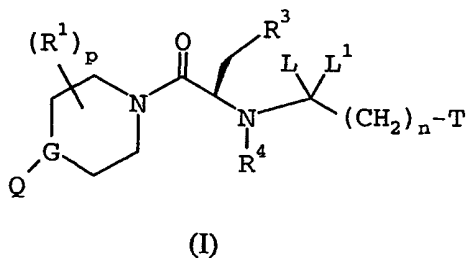
; and

- 10 h) coupling the compound of formula 9 with a compound having a



formula, to afford a compound of formula I.

Yet another aspect of the present invention is a process for preparing a compound of formula I:

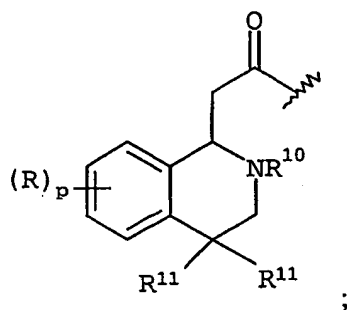


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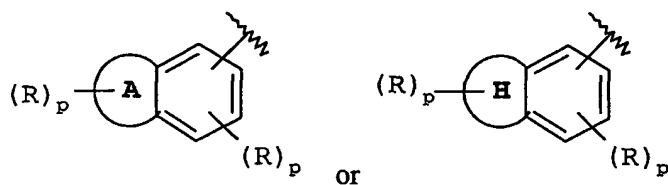
or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

-45-

5 -CLL¹-(CH₂)_n-T is:



Q represents a moiety:



10 G is CR¹ or N;



is a 5- or 6-membered carbocyclyl wherein the carbocyclyl being saturated, partially saturated or aromatic ring, and the carbocyclyl being optionally substituted with one to three substituents independently selected from R;

15



is a 5- or 6-membered heterocyclyl,

wherein the heterocyclyl being saturated, partially saturated or aromatic ring, provided that when G is carbon, the 5-membered nitrogen containing heterocyclyl does not contain a carbon-carbon double bond in the ring; and

20 wherein the heterocyclyl being optionally substituted with one to three substituents independently selected from R²;

-46-

- 5 R is independently:
hydrogen,
hydroxy,
(D)cyano,
halo,
10 C₁-C₈ alkyl,
C₁-C₈ alkoxy,
C₃-C₇ cycloalkyl,
C₁-C₄ haloalkyl,
(D)heterocyclyl
15 (D)C(O)R⁸,
(D)C(O)(CH₂)_nN(R⁸)₂,
C₁-C₈ alkyl-N(R⁸)₂,
(D)OR⁸,
(D)OCOR⁸,
20 (D)OC(O)N(R⁸)₂,
(D)N(R⁸)₂,
(D)NR⁸C(O)R⁸,
(D)NR⁸C(O)OR⁸,
(D)NR⁸C(O)N(R⁸)₂,
25 (D)NR⁸SO₂R⁸,
(D)SR⁸,
(D)SOR⁸,
(D)SO₂R⁸, or
(D)SO₂N(R⁸)₂;
30 R¹ is independently:

-47-

- 5 hydrogen, CONH(C₁-C₈ alkyl), C₁-C₈ alkyl, (D)phenyl, (D)C₃-C₇ cycloalkyl or oxo, provided that oxo is not attached to the same carbon that attached to nitrogen which forms an amide bond when G is N;

R² is independently:

- 10 hydrogen,
C₁-C₈ alkyl,
C₃-C₇ cycloalkyl,
C(O)NH₂,
C(O)O(C₁-C₆ alkyl),
15 C(O)(CH₂)_nN(C₁-C₆ alkyl)₂,
C(O)-phenyl,
SO₂(C₁-C₆ alkyl),
SO₂N(C₁-C₆ alkyl)₂,
SO₂-phenyl or
20 C(O)(C₁-C₆ alkyl);

R³ is independently: aryl or thienyl;

wherein aryl and thienyl are optionally substituted with one to three substituents selected from the group consisting of:

- 25 cyano, halo, C₁-C₈ alkyl, (D)C₃-C₇ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkyl and C₁-C₄ haloalkyloxy;

-48-

5 R^4 is independently:

hydrogen, C_1 - C_8 alkyl, $C(O)R^8$, $C(O)OR^8$, C_3 - C_7 cycloalkyl or $(CH_2)_nO(C_1$ - C_8 alkyl), wherein n is 2-8;

each R^8 is independently:

10 hydrogen,
 aryl, wherein aryl being phenyl or naphthyl,
 C_1 - C_8 alkyl,
 C_1 - C_4 alkoxy, or
 $(CH_2)_nC_1$ - C_4 haloalkyl, wherein n is 2-8;

15 each R^{10} is independently:

hydrogen, $(C_1$ - $C_8)$ alkyl, $C(O)C_1$ - C_8 alkyl, aryl, C_3 - C_7 cycloalkyl, or protecting group selected from Boc, CBZ or FMOC;

each R^{11} is independently: hydrogen or C_1 - C_8 alkyl;

20

D is a bond or $-(CH_2)_n$;

n is 0-8;

p is 0-4; and

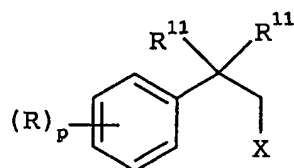
q is 0-1;

25

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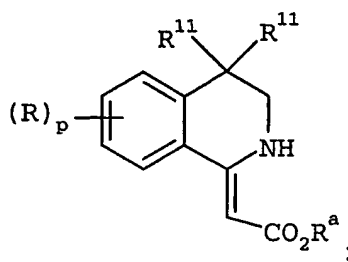
5 comprising the steps of:

a) reacting a compound formula 1:



(1)

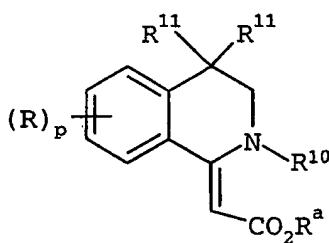
10 wherein X is halo, and R^{11} is independently, hydrogen or C1-C4 alkyl, with $CNCH_2CO_2R^a$ wherein R^a is C₁-C₈ alkyl or benzyl to afford a compound of formula 2:



(2)

15

b) protecting the compound of formula 2 to form the compound of formula 3:

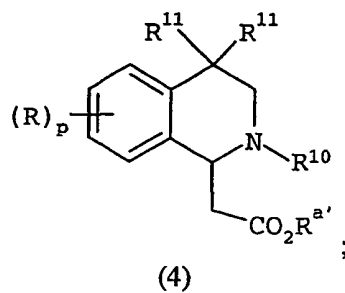


(3)

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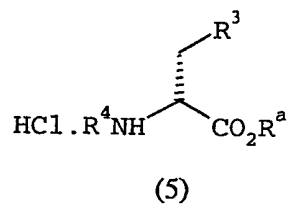
-50-

- 5 c) hydrogenating the compound of formula 3 to afford a compound of formula 4:

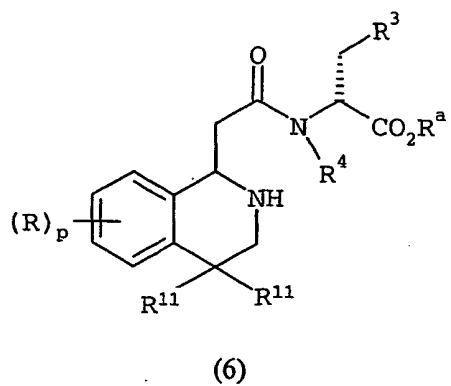


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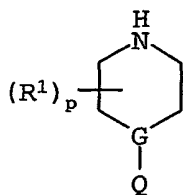
- d) coupling the compound of formula 4 wherein R^{a'} is hydrogen or R^a, with a compound of formula 5,



- 15 to afford a compound of formula 6,



- e) coupling the compound of formula 6 with a compound having a



- 20 formula, to afford a compound of formula I.

5

Throughout the instant application, the following terms have the indicated meanings:

The term "5- or 6- membered carbocycl" refers to carbocycl ring that is saturated, partially saturated or aromatic ring. The saturated carbocycl ring includes 5-
10 or 6- membered cycloalkyl, which may be optionally substituted with other hydrocarbon substituents. Examples of cycloalkyl are cyclopentyl and cyclohexyl, and cycloalkyl as defined above may be optionally substituted with designated number of substituents as set forth in the embodiment recited above.

The partially saturated carbocycl ring includes 5- or -6 membered
15 cycloalkenyl, which may be optionally substituted with other hydrocarbon substituents. Examples of cycloalkenyl are cyclopentenyl and cyclohexenyl, and cycloalkenyl as defined above may be optionally substituted with designated number of substituents as set forth in the embodiment recited above.

The aromatic carbocycl ring refers to 5- or 6-membered aromatic ring
20 such as phenyl.

The "5- or 6-membered heterocycl" refers to a heterocyclic ring that is saturated, partially saturated or aromatic ring (i.e., heteroaryl) containing from one to four heteroatoms selected from N, O or S. The "heterocycl" includes "nitrogen containing heterocycl," which contains from one to four nitrogen atoms and optionally further
25 contains one other heteroatom selected from O or S. Heterocycl as defined above may be optionally substituted with designated number of substituents as set forth in the embodiment recited above. The examples of 5- or 6-membered heterocycl includes, but are not limited to: piperidinyl, pyrrolidinyl, pyrrolinyl, isoxazolyl, oxazolyl, thiazolyl, triazolyl, tetrazolyl, thiadiazolyl, or oxadiazolyl, furanyl, thienyl, thiazolyl, isoxazolyl,
30 imidazolyl, oxazolyl, isoxazolyl, pyrazoyl, pyrrolyl, pyrazinyl, pyridyl, pyrimidyl, and purinyl and the like.

The term "alkyl," unless otherwise indicated, refers to those alkyl groups of a designated number of carbon atoms of either a straight or branched saturated configuration. Examples of "alkyl" includes, but are not limited to methyl, ethyl, n-
35 propyl, isopropyl, n-butyl, isobutyl, sec-butyl and t-butyl, pentyl, hexyl, neopenyl,

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5 isopentyl and the like. Alkyl as defined above may be optionally substituted with a designated number of substituents as set forth in the embodiment recited above.

The term "alkenyl" means hydrocarbon chain of a specified number of carbon atoms of either a straight or branched configuration and having at least one carbon-carbon double bond, which may occur at any point along the chain, such as
10 ethenyl, propenyl, butenyl, pentenyl, vinyl, alkyl, 2-butenyl and the like. Alkenyl as defined above may be optionally substituted with designated number of substituents as set forth in the embodiment recited above.

The term "haloalkyl" is an alkyl group of indicated number of carbon atoms, which is substituted with one to five halo atoms selected from F, Br, Cl and I. An
15 example of a haloalkyl group is trifluoromethyl.

The term "alkoxy" represents an alkyl group of indicated number of carbon atoms attached through an oxygen bridge, such as methoxy, ethoxy, propoxy, isopropoxy, butoxy, tert-butoxy, pentoxy, and the like. Alkoxy as defined above may be optionally substituted with a designated number of substituents as set forth in the
20 embodiment recited above.

The term "cycloalkyl" refers to a ring composed of 3 to 7 methylene groups, each of which may be optionally substituted with other hydrocarbon substituents. Examples of cycloalkyl includes, but are not limited to: cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl, and the like. Cycloalkyl as defined above may
25 be optionally substituted with a designated number of substituents as set forth in the embodiment recited above.

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5 The term "halo" refers to fluoro, chloro, bromo and iodo.

 The term "haloalkyloxy" represents a haloalkyl group of indicated number of carbon atoms attached through an oxygen bridge, such as OCF_3 . "Haloalkyloxy" as defined above may be optionally substituted with a designated number of substituents as set forth in the embodiment recited above.

10 The term "aryl" refers to phenyl, naphthyl, anthracenyl, phenanthrenyl and the like which is optionally substituted with a designated number of substituents as set forth in the embodiment recited above.

 The term "heteroaryl" refers to monocyclic or bicyclic aromatic ring of 5- to 10-carbon atoms containing from one to four heteroatoms selected from O, N, or S, and the heteroaryl being optionally substituted with a designated number of substituents as set forth in the embodiment recited above. Examples of heteroaryl are, but are not limited to furanyl, thienyl, thiazolyl, imidazolyl, isoxazolyl, oxazolyl, pyrazolyl, pyrrolyl, pyrazinyl, pyridyl, pyrimidyl, and purinyl, cinnolinyl, benzothienyl, benzotriazolyl, benzoxazolyl, quinoline, isoquinoline and the like.

20 The "heterocyclyl" is defined as a monocyclic, bicyclic, or tricyclic ring of 5 to 14 carbon atoms which are saturated or partially saturated containing from one to four heteroatoms selected from N, O or S. The "heterocyclyl" includes "nitrogen containing heterocyclyl," which contains from one to four nitrogen atoms and optionally further contains one other heteroatom selected from O or S. Heterocyclyl as defined above may be optionally substituted with a designated number of substituents as set forth in the embodiment recited above.

 A mammal as used in here includes a human and a warm-blooded animal such as a cat, a dog and the like.

30 The term "composition" or "formulation", as in pharmaceutical composition or formulation, is intended to encompass a product comprising the active ingredient(s), and the inert ingredient(s) that make up the carrier. Accordingly, the pharmaceutical compositions of the present invention encompass any composition made by admixing a compound of the present invention (a compound of formula I) and a pharmaceutically acceptable carrier.

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5 The term "pharmaceutical" when used herein as an adjective means substantially non-deleterious to the recipient mammal.

 The term "unit dosage form" refers to physically discrete units suitable as unitary dosages for human subjects and other non-human animals such as warm-blooded animals each unit containing a predetermined quantity of active ingredient (a compound
10 of formula I) calculated to produce the desired therapeutic effect in association with a suitable pharmaceutical carrier.

 The term "treating" or "preventing" as used herein includes its generally accepted meanings, *i.e.*, preventing, prohibiting, restraining, alleviating, ameliorating, slowing, stopping, or reversing the progression or severity of a pathological condition, or
15 sequela thereof as described herein.

 "Erectile dysfunction" is a disorder involving the failure of a male mammal to achieve erection, ejaculation, or both. Symptoms of erectile dysfunction include an inability to achieve or maintain an erection, ejaculatory failure, premature ejaculation, and the inability to achieve an orgasm. An increase in erectile dysfunction is
20 often associated with age and is generally caused by a physical disease or as a side effect of drug treatment.

 "Female sexual dysfunction" encompasses, without limitation, conditions such as a lack of sexual desire and related arousal disorders, inhibited orgasm, lubrication difficulties, and vaginismus.

25 Because certain compounds of the invention contain an acidic moiety (e.g., carboxy), the compound of formula I may exist as a pharmaceutical base addition salt thereof. Such salts include those derived from inorganic bases such as ammonium and alkali and alkaline earth metal hydroxides, carbonates, bicarbonates and the like, as well as salts derived from basic organic amines such as aliphatic and aromatic amines,
30 aliphatic diamines, hydroxy alkamines, and the like.

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5 Because certain compounds of the invention contain a basic moiety (e.g., amino), the compound of formula I can also exist as a pharmaceutical acid addition salt. Such salts include sulfate, pyrosulfate, bisulfate, sulfite, bisulfite, phosphate, mono-hydrogenphosphate, dihydrogenphosphate, metaphosphate, pyrophosphate, chloride, bromide, iodide, acetate, propionate, decanoate, caprylate, acrylate, formate, isobutyrate, 10 heptanoate, propiolate, oxalate, malonate, succinate, suberate, sebacate, fumarate, maleate, 2-butyne-1,4 dioate, 3-hexyne-2, 5-dioate, benzoate, chlorobenzoate, hydroxybenzoate, methoxybenzoate, phthalate, xylenesulfonate, phenylacetate, phenylpropionate, phenylbutyrate, citrate, lactate, hippurate, beta-hydroxybutyrate, glycollate, maleate, tartrate, methanesulfonate, propanesulfonate, naphthalene-1-sulfonate, 15 naphthalene-2-sulfonate, mandelate and the like salts. Preferred salt form of compound of formula I is an acid addition salts, more specifically hydrochloride salt.

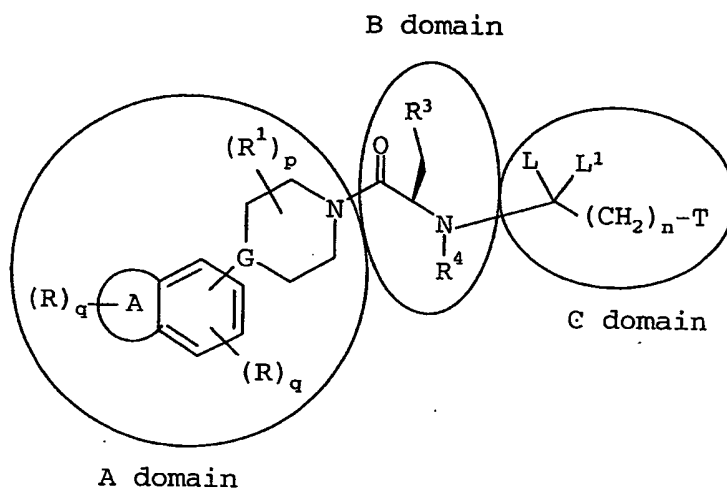
 Some of the compounds described herein may exist as tautomers such as keto-enol tautomers. The individual tautomers as well as mixtures thereof are encompassed within the scope of the present invention.

20 Utility

 Compounds of formula I are effective as melanocortin receptor modulators, particularly as agonists of the human MC-4 receptor. As melanocortin receptor agonists, the compounds of formula I are useful in the treatment of diseases, disorders or conditions responsive to the activation of one or more of the melanocortin 25 receptors including, but not limited to, MC-1, MC-2, MC-3, MC-4, and MC-5. Diseases, disorders or conditions receptive to treatment with a MC-4 agonist include those mentioned above and those described in WO 00/74679, the teachings of which are herein incorporated by reference. In particular diseases, disorders or conditions receptive to treatment with a MC-4 agonist include obesity or diabetes mellitus, male or female sexual 30 dysfunction, more specifically erectile dysfunction.

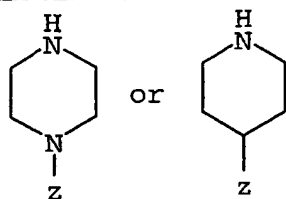
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- 5 When describing various aspects of the present compounds of formula I, the terms "A domain", "B domain" and "C domain" are used below. This domain concept is illustrated below:



- 10 The following listing provides some of examples "A domain", "B domain" and "C domain" of the compound of formula I. These listings are provided as illustrative purposes and as such are not meant to be limiting.

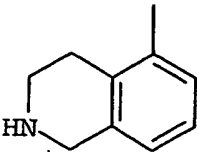
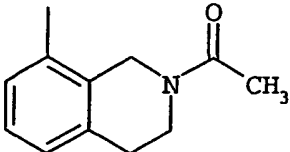
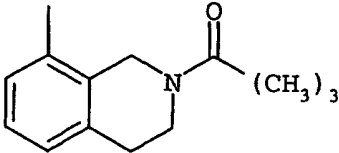
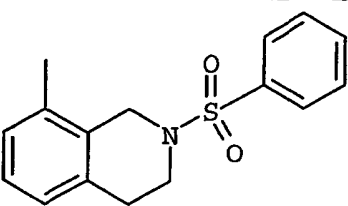
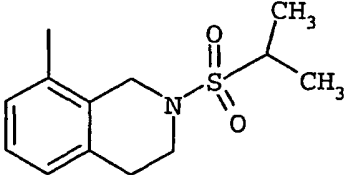
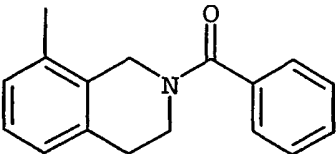
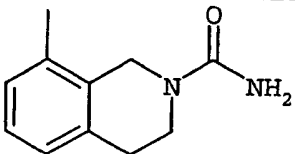
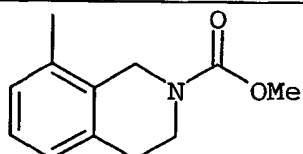
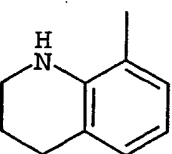
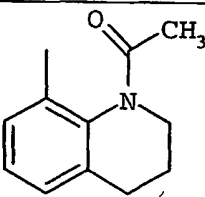
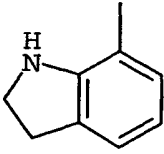
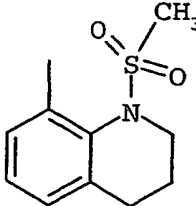
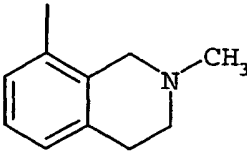
-57-

5 Examples of "A domain":

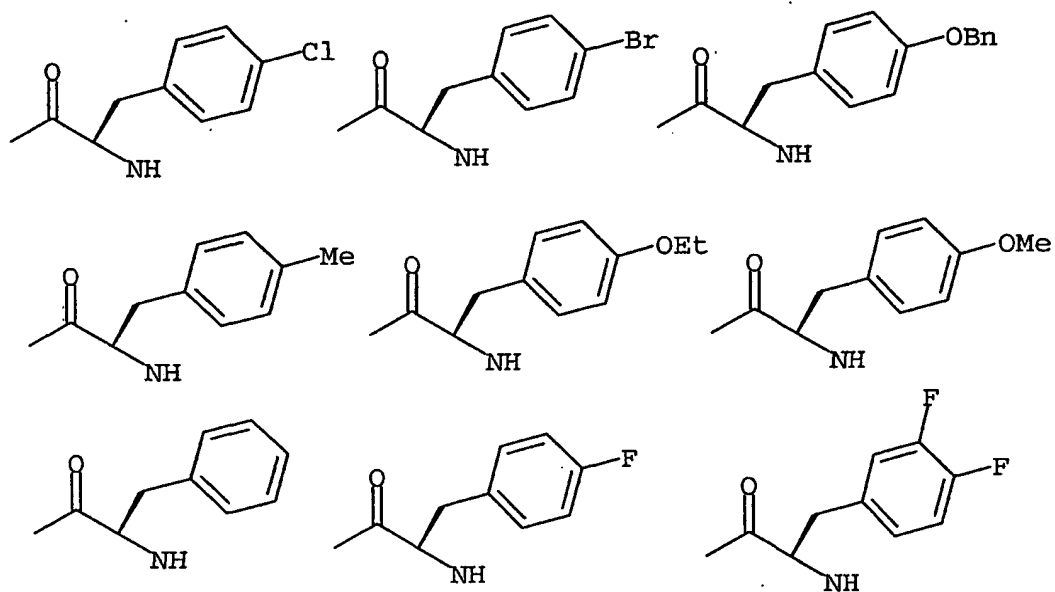
where Z is shown in the table below:

Z	Z

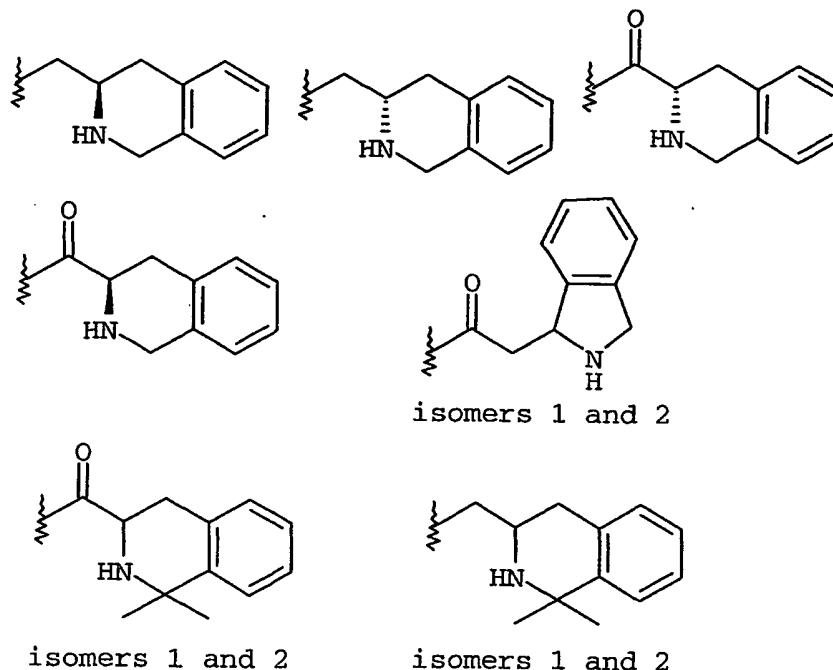
-58-

Z	Z
	
	
	
	
	
	
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5

Examples of "B domain":

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5 Examples of C domain:Formulation

10 The compound of formula I is preferably formulated in a unit dosage form prior to administration. Accordingly the present invention also includes a pharmaceutical composition comprising a compound of formula I and a suitable pharmaceutical carrier.

The present pharmaceutical compositions are prepared by known procedures using well-known and readily available ingredients. In making the formulations of the present invention, the active ingredient (a compound of formula I) is usually mixed with a carrier, or diluted by a carrier, or enclosed within a carrier, which may be in the form of a capsule, sachet, paper or other container. When the carrier serves as a diluent, it may be a solid, semisolid or liquid material which acts as a vehicle, excipient or medium for the active ingredient. Thus, the compositions can be in the form of tablets, pills, powders, lozenges, sachets, cachets, elixirs, suspensions, emulsions, solutions, syrups, aerosol (as a solid or in a liquid medium), soft and hard gelatin capsules, suppositories, sterile injectable solutions and sterile packaged powders.

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5 Some examples of suitable carriers, excipients, and diluents include
lactose, dextrose, sucrose, sorbitol, mannitol, starches, gum acacia, calcium phosphate,
alginate, tragacanth, gelatin, calcium silicate, microcrystalline cellulose,
polyvinylpyrrolidone, cellulose, water syrup, methyl cellulose, methyl and
propylhydroxybenzoates, talc, magnesium stearate and mineral oil. The formulations can
10 additionally include lubricating agents, wetting agents, emulsifying and suspending
agents, preserving agents, sweetening agents or flavoring agents. The compositions of
the invention may be formulated so as to provide quick, sustained or delayed release of
the active ingredient after administration to the patient.

15 Dosage:

The specific dose administered is determined by the particular
circumstances surrounding each situation. These circumstances include, the route of
administration, the prior medical history of the recipient, the pathological condition or
symptom being treated, the severity of the condition/symptom being treated, and the age
20 and sex of the recipient. Additionally, it would be understood that the therapeutic dosage
administered can be determined by the physician in the light of the relevant
circumstances.

Generally, an effective minimum daily dose of a compound of formula I is
about 1, 5, 10, 15, or 20 mg. Typically, an effective maximum dose is about 500, 100,
25 60, 50, or 40 mg. The suitable dose may be determined in accordance with the standard
practice in the medical arts of "dose titrating" the recipient, which involves administering
a low dose of the compound initially and then gradually increasing the dose until the
desired therapeutic effect is observed.

30

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5 Route of Administration

The compounds may be administered by a variety of routes including the oral, rectal, transdermal, subcutaneous, topical, intravenous, intramuscular or intranasal routes.

10 Combination Therapy

Compounds of formula I may be used in combination with other drugs that are used in the treatment of the diseases or conditions for which compounds of formula I are useful. Such other drugs may be administered by a route and in an amount commonly used therefore, contemporaneously or sequentially with a compound of formula I. When
15 a compound of formula I is used contemporaneously with one or more other drugs, a pharmaceutical composition containing such other drugs in addition to the compound of formula I is preferred. Accordingly, the pharmaceutical compositions of the present invention include those that also contain one or more other active ingredients in addition to a compound of formula I. Examples of other active ingredients that may be combined
20 with a compound of formula I, either administered separately or in the same pharmaceutical compositions, include but are not limited to:

- (a) insulin sensitizers including (i) PPAR-gamma agonists such as the glitazones (e.g., troglitazone, pioglitazone, englitazone, MCC-555, BRL49653 and the like) and compounds disclosed in WO97/27857, WO 97/28115, WO 97/28137 and WO97/27847;
- 25 (ii) biguanides such as metformin and phenformin;
- (b) insulin or insulin mimetics;
- (c) sulfonylureas such as tolbutamide and glipizide;
- (d) alpha-glucosidase inhibitors (such as acarbose),
- (e) cholesterol lowering agents such as (i) HMG-CoA reductase inhibitors
30 (lovastatin, simvastatin and pravastatin, fluvastatin, atorvastatin, and other statins), (ii) sequestrants (cholestyramine, colestipol and a dialkylaminoalkyl derivatives of a cross-linked dextran), (iii) nicotiny alcohol nicotinic acid or a salt thereof, (iv) proliferator-activater receptor alpha-agonists such as fenofibric acid derivatives (gemfibrozil, clofibrat, fenofibrate and benzaifibrate), (v) inhibitors of cholesterol absorption such as

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- 5 beta-sitosterol and acyl CoA:cholesterol acyltransferase inhibitors such as melinamide,
 (vi) probucol, (vii) vitamin E, and (viii) thyromimetics;
 (f) PPAR-delta agonists such as those disclosed in WO97/28149;
 (g) antiobesity compounds such as fenfluramine, dexfenfluramine, phentermine,
 sibutramine, orlistat, and beta-3 adrenergic receptor agonists;
 10 (h) feeding behavior modifying agents such as neuropeptide Y antagonists (e.g.,
 neuropeptide Y5) as disclosed in WO 97/19682, WO 97/20820, WO 97/20821, WO
 97/20822 and WO 97/20823;
 (i) PPAR-alpha agonists as described in WO 97/36579;
 (j) PPAR-gamma antagonists as described in WO97/10813;
 15 (k) serotonin reuptake inhibitors, such as fluoxetine, sertraline and 5HT2C agonists;
 (l) growth hormone secretagogues such as MK-0677 and ghrelin antagonists; and
 (m) agents useful in the treatment of male and/or female sexual dysfunction, such as
 phosphodiester V inhibitors including sildenafil and ICI-351, and alpha-2 adrenergic
 receptor antagonists including phentolamine mesylate; and dopamine-receptor agonists
 20 such as apomorphine, and MC3R, MCH, His3, leptin and opioids.

Biological Assays

A. Binding Assay:

- The radioligand binding assay is used to identify competitive inhibitors of
 25 ¹²⁵I-NDP-alpha-MSH binding to cloned human MCRs using membranes from stably
 transfected human embryonic kidney (HEK) 293 cells.

- HEK 293 cells transfected with human or rat melanocortin receptors are grown
 either as adherent monolayers or suspension culture. Monolayer cells are grown in roller
 bottle cultures at 37°C and 5%CO₂/air atmosphere in a 3:1 mixture of Dulbecco's
 30 modified Eagle medium (DMEM) and Ham's F12 containing 25 mM L-glucose, 100
 units/ml penicillin G, 100 microgram/ml streptomycin, 250 nanogram/ml amphotericin B,
 300 microgram/ml gentamicin and supplemented with 5% fetal bovine serum. Monolayer
 cells are adapted to suspension culture (Berg et al., *Biotechniques* Vol. 14, No.6, 1993)
 and are grown in either spinner or shaker flasks (37°C and 7.5% CO₂/air overlay) in a
 35 modified DME/F12 medium containing 0.1 mM CaCl₂, 2% equine serum and 100

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5 microgram/ml sodium heparin to prevent cell-cell aggregation. Cells are harvested by centrifugation, washed in PBS, and pellets are stored frozen at -80°C until membrane preparations.

The cell pellets are resuspended in 10 volumes of membrane preparation buffer (*i.e.*, 1g pellet to 10 ml buffer) having the following composition: 50 mM Tris pH
10 7.5 @ 4°C, 250mM sucrose, 1mM MgCl₂, Complete® EDTA-free protease inhibitor tablet (Boehringer Mannheim), and 24 micrograms/ml DNase I (Sigma, St. Louis, MO). The cells are homogenized with a motor-driven dounce using 20 strokes, and the homogenate is centrifuged at 38,000 x g at 4°C for 40 minutes. The pellets are resuspended in membrane preparation buffer at a concentration of 2.5-7.5 mg/ml and 1
15 milliliter aliquots of membrane homogenates are quickly frozen in liquid nitrogen and then stored at -80°C.

Solutions of a compound of formula I (300 picomolar to 30 micromolar) or unlabelled NDP-alpha-MSH (1 picomolar to 100 nanomolar) are added to 150 microliters of membrane binding buffer to yield final concentrations (listed in parentheses). The
20 membrane binding buffer has the following composition: 25 mM HEPES pH 7.5; 10 mM CaCl₂; 0.3% BSA). One hundred fifty microliters of membrane binding buffer containing 0.5-5.0 microgram membrane protein is added, followed by 50 nanomolar ¹²⁵I-NDP-alpha-MSH to final concentration of 100 picomolar. Additionally, fifty microliters of SPA beads (5 mg/ml) are added and the resulting mixture is agitated briefly
25 and incubated for 10 hours at r.t. The radioactivity is quantified in a Wallace Trilux Microplate Scintillation counter. IC₅₀ values obtained in competition assays are converted to affinity constants (K_i values) using the Cheng-Prusoff equation: $K_i = IC_{50} / (1 + D/K_d)$.

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5 B. Functional Assay:

Functional cell based assays are developed to discriminate agonists and antagonists.

Agonist Assay: HEK 293 cells stably expressing a human melanocortin receptor (see *e.g.*, Yang, et al., *Mol-Endocrinol.*, 11(3): 274-80, 1997) are dissociated
10 from tissue culture flasks using a trypsin/EDTA solution(0.25%; Life Technologies, Rockville, MD). Cells are collected by centrifugation and resuspended in DMEM (Life Technologies, Rockville, MD) supplemented with 1% L-glutamine and 0.5% fetal bovine serum. Cells are counted and diluted to 4.5×10^5 /ml.

A compound of formula I is diluted in DMSO
15 (3×10^{-5} to 3×10^{-10} M final concentrations) and 0.05 volume of compound solution is added to 0.95 volumes of cell suspension; the final DMSO concentration is 0.5%. After incubation at 37°C/5% CO₂ for 5 hours, cells are lysed by addition of luciferin solution (50mM Tris, 1mM MgCl₂, 0.2% Triton-X100, 5mM DTT, 500 micromolar Coenzyme A, 150 micromolar ATP, and 440 micromolar luciferin) to quantify the activity of the
20 reporter gene luciferase, an indirect measurement of intracellular cAMP production.

Luciferase activity is measured from the cell lysate using a Wallace Victor 2 luminometer. The amount of lumen production which results from a compound of formula I is compared to that amount of lumens produced in response to NDP-alpha-MSH, defined as a 100% agonist, to obtain the relative efficacy of a compound. The
25 EC₅₀ is defined as the compound concentration that results in half maximal stimulation, when compared to its own maximal level of stimulation.

Antagonist assay: Antagonist activity is defined as the ability of a compound to block lumen production in response to NDP-alpha-MSH. Concentration-response curves are generated for NDP-alpha-MSH in the absence and presence of a fixed
30 concentration of a solution of a compound of formula I ($10 \times K_i$ from binding assays). Suspensions of MCR-expressing cells are prepared and are incubated with NDP-alpha-MSH and compound solutions for 5 hours as described above. The assay is terminated by the addition of luciferin reagent and lumen production is quantified. Antagonist potency is determined from the rightward shift of the EC₅₀ value in the absence of a compound of
35 formula I using the equation: $K_b = \text{Concentration of Antagonist}/[(EC_{50}'/EC_{50})-1]$.

5

Whole Cell cAMP Accumulation AssayCompound Preparation

In the agonist assay, compounds are prepared as 10mM and NDP-alpha-MSH (control) as 33.3 μ M stock solutions in 100 % DMSO. These are serially diluted in 100 % DMSO. The compound plate is further diluted 1:200 in compound dilution buffer (HBSS-092, 1 mM Ascorbic Acid, 1 mM IBMX, 0.6% DMSO, 0.1% BSA). The final concentration range being 10 μ M–100 pM for compound and 33.33 nM–0.3 pM for control in 0.5 % DMSO. Transfer 20 μ l from this plate into four PET 96-well plates (all assays are performed in duplicate for each receptor).

15 Cell Culture and Cell Stimulation

HEK 293 cells stably transfected with the MC3R and MC4R were grown in DMEM containing 10% FBS and 1% Antibiotic/Antimycotic Solution. On the day of the assay the cells were dislodged with enzyme free cell dissociation solution and resuspended in cell buffer (HBSS-092, 0.1% BSA, 10 mM HEPES) at 1×10^6 cells/ml.

20 Add 40 μ l of cells/well to the PET 96-well plates containing 20 microliter diluted compound and control. Incubate @ 37°C in a water bath for 20 minutes. Stop the assay by adding 50 μ l Quench Buffer (50 mM Na Acetate, 0.25% Triton X-100).

Radioligand Binding Assays

Radioligand binding assays were run in SPA buffer (50mM Sodium Acetate, 0.1% BSA). The beads, antibody and radioligand were diluted in SPA buffer to provide sufficient volume for each 96-well plate. To each quenched assay well was added 100 microliter cocktail containing 33.33 microliter of beads, 33.33 microliter antibody and 33.33 microliter 125 I-cAMP. This was based on a final concentration of 6.3 mg/ml beads, 0.65 % anti-goat antibody and 61 pM of 125 I-cAMP (containing 25000-30000 CPM) in a final assay volume of 210 microliter. The plates were counted in a Wallac MicroBeta counter after a 12-hour incubation.

The data was converted to pmoles cAMP using a standard curve assayed under the same conditions. The data was analyzed using Activity Base software to generate agonist potencies (EC50) and percent relative efficacy data to NDP-alpha-MSH.

35 C. In vivo Food Intake Models:

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5 1) Overnight food intake. Sprague Dawley rats are injected
intracerebroventricularly with a test compound in 400 nL of 50% propylene
glyco/artificial cerebrospinal fluid one hour prior to onset of dark cycle (12 hours). Food
intake is determined using a computerized system in which each rat's food is placed on a
computer monitored balance. Cumulative food intake for 16 hours post compound
10 administration is measured.

 2) Food intake in diet induced obese mice. Male C57/B16J mice
maintained on a high fat diet (60% fat calories) for 6.5 months from 4 weeks of age are
dosed intraperitoneally with a compound of formula I. Food intake and body weight are
measured over an eight-day period. Biochemical parameters relating to obesity, including
15 leptin, insulin, triglyceride, free fatty acid, cholesterol and serum glucose levels are
determined.

D. Rat Ex Copula Assay:

 Sexually mature male Caesarian Derived Sprague Dawley (CD) rats (over
20 60 days old) are used with the suspensory ligament surgically removed to prevent
retraction of the penis back into the penile sheath during the ex copula evaluations.
Animals receive food and water *ad lib* and are kept on a normal light/dark cycle. Studies
are conducted during the light cycle.

 1) Conditioning to Supine Restraint for Ex Copula Reflex Tests. This
25 conditioning takes about 4 days. Day 1, the animals are placed in a darkened restrainer
and left for 15-30 minutes. Day 2, the animals are restrained in a supine position in the
restrainer for 15-30 minutes. Day 3, the animals are restrained in the supine position with
the penile sheath retracted for 15-30 minutes. Day 4, the animals are restrained in the
supine position with the penile sheath retracted until penile responses are observed. Some
30 animals require additional days of conditioning before they are completely acclimated to
the procedures; non-responders are removed from further evaluation. After any handling
or evaluation, animals are given a treat to ensure positive reinforcement.

 2) Ex Copula Reflex Tests. Rats are gently restrained in a supine position
with their anterior torso placed inside a cylinder of adequate size to allow for normal head
35 and paw grooming. For a 400-500 gram rat, the diameter of the cylinder is approximately

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5 8 cm. The lower torso and hind limbs are restrained with a non-adhesive material (vetrap). An additional piece of vetrap with a hole in it, through which the glans penis will be passed, is fastened over the animal to maintain the perpetual sheath in a retracted position. Penile responses will be observed, typically termed *ex copulu* genital reflex tests. Typically, a series of penile erections will occur spontaneously within a few
10 minutes after sheath retraction. The types of normal reflexogenic erectile responses include elongation, engorgement, cup and flip. An elongation is classified as an extension of the penile body. Engorgement is a dilation of the glans penis. A cup is defined as an intense erection where the distal margin of the glans penis momentarily flares open to form a cup. A flip is a dorsiflexion of the penile body.

15 Baseline and/or vehicle evaluations are conducted to determine how and if an animal will respond. Some animals have a long duration until the first response while others are non-responders altogether. During this baseline evaluation, latency to first response time, number and type of responses are recorded. The testing time frame is 15 minutes after the first response.

20 After a minimum of 1 day between evaluations, these same animals are administered a compound of formula I at 20 mg/kg and evaluated for penile reflexes. All evaluations are videotaped and scored later. Data are collected and analyzed using paired 2 tailed t-tests to compared baseline and/or vehicle evaluations to drug treated evaluations for individual animals. Groups of a minimum of 4 animals are utilized to reduce
25 variability.

 Positive reference controls are included in each study to assure the validity of the study. Animals can be dosed by a number of routes of administration depending on the nature of the study to be performed. The routes of administration includes intravenous (IV), intraperitoneal (IP), subcutaneous (SC) and intracerebral ventricular
30 (ICY).

E. Models of Female Sexual Dysfunction:

 Rodent assays relevant to female sexual receptivity include the behavioral model of lordosis and direct observations of copulatory activity. There is also a urethrogenital reflex model in anesthetized spinally transected rats for measuring orgasm
35 in both male and female rats. These and other established animal models of female

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- 5 sexual dysfunction are described in McKenna, et al., *Am. J. Physiol.*, (Regulatory Integrative Comp. Physiol 30):R1276-R1285, 1991; McKenna, et al., *Pharm. Bioch. Behav.*, 40:151-156, 1991; and Takahashi, et al., *Brain Res.*, 359:194-207, 1985.

Preparation of the Compounds of the Invention

- 10 Preparation of the compounds of the present invention may be carried out via sequential or convergent synthetic routes. The skilled artisan will recognize that, in general, the three domains of a compound of formula I are connected via amide bonds. The B and C domains are optionally connected via a reduced or partially reduced amide bond (*e.g.*, via reductive amination). The skilled artisan can, therefore, readily envision
15 numerous routes and methods of connecting the three domains via standard peptide coupling reaction conditions.

- The phrase "standard peptide coupling reaction conditions" means coupling a carboxylic acid with an amine using an acid activating agent such as EDC, dicyclohexylcarbodiimide and benzotriazol-1-yloxytris(dimethylamino)phosphonium
20 hexafluorophosphate in a inert solvent such as DCM in the presence of a catalyst such as HOBT. The uses of protective groups for amine and carboxylic acids to facilitate the desired reaction and minimize undesired reactions are well documented. Conditions required to remove protecting groups which may be present can be found in Greene, *et al.*, *Protective Groups in Organic Synthesis*, John Wiley & Sons, Inc., New York, NY
25 1991.

- CBZ, Boc and Fmoc protecting groups are used extensively in the synthesis, and their removal conditions are well known to those skilled in the art. For example, removal of CBZ groups can be achieved by catalytic hydrogenation with hydrogen in the presence of a metal such as palladium on activated carbon in a protic
30 solvent such as ethanol. In cases where catalytic hydrogenation is contraindicated by the presence of other potentially reactive functionality, removal of CBZ can also be achieved by treatment with a solution of hydrogen bromide in acetic acid. Removal of Boc protecting groups is carried out in a solvent such as methylene chloride, methanol or ethyl acetate with a strong acid, such as TFA or HCl or hydrogen chloride gas. Fmoc
35 protecting groups may be removed by 20% piperidine in DMF.

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5 The compounds of formula I, when they exist as a diastereomeric mixture, may be separated into diastereomeric pairs of enantiomers by fractional crystallization from a suitable solvent such as methanol, ethyl acetate or a mixture thereof. The pair of enantiomers thus obtained may be separated into individual stereoisomers by conventional means by using an optically active acid as a resolving agent. Alternatively, 10 any enantiomer of a compound of the formula I may be obtained by stereospecific synthesis using optically pure starting materials or reagents of known configuration.

 The compounds of the present invention can be prepared according to the procedure of the following schemes and examples, which may further illustrate details for the preparation of the compounds of the present invention. The compounds illustrated in 15 the examples are, however, not to be construed as forming the only genus that is considered as the present invention.

 In the Schemes, Preparations and Examples below, various reagent symbols and abbreviations have the following meanings:

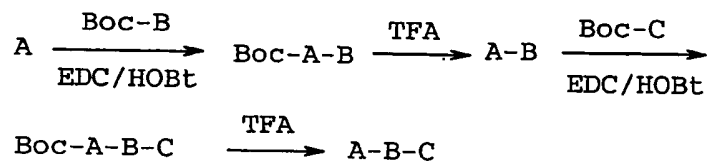
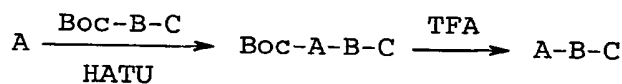
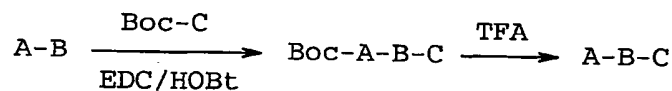
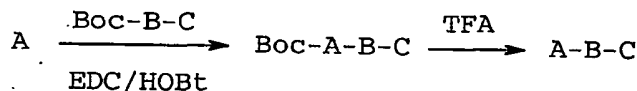
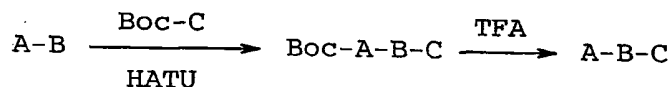
	BINAP	2,2'-Bis(diphenylphosphino)-1,1'-binaphthyl
20	Boc	t-butoxycarbonyl
	CBZ	benzyloxycarbonyl
	DCM	dichloromethane
	DEAD	diethyl azodicarboxylate
	DIAD	diisopropyl azodicarboxylate
25	DIPEA	diisopropylethylamine
	DMAP	4-dimethylamino pyridine
	DMF	N,N-dimethylformamide
	DMSO	dimethylsulfoxide
	eq.	equivalent(s)
30	EDC	1-(3-dimethylaminopropyl)-3-ethylcarbodiimide HCl
	ESI-MS	electron spray ion-mass spectroscopy
	Et	ethyl
	EtOAc	ethyl acetate
	Fmoc	9-fluorenylmethyl carbamate

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5	HATU	O-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate
	HOAT:	1-hydroxy-7-azabenzotriazole
	HOBT	1-hydroxybenzotriazole hydrate
	HPLC	high performance liquid chromatography
10	HRMS	high resolution mass spectroscopy
	h (hr)	hour(s)
	LRMS	low resolution mass spectroscopy
	Me	methyl
	Ms	methanesulfonyl
15	NMM	4-methylmorpholine
	Pd ₂ (dba) ₃	tris(dibenzylideneacetone) dipalladium(0)
	Ph	phenyl
	Phe	phenylalanine
	Pr	propyl
20	r.t.	room temperature
	TBAF	tetrabutylammonium fluoride
	TBS	tertbutyldimethylsilyl
	TFA	trifluoroacetic acid
	TEA	triethylamine
25	THF	tetrahydrofuran
	Tic	1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid
	TLC	thin-layer chromatography

Reaction Scheme 1: Coupling procedures

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Procedure 1**Procedure 2****Procedure 3****Procedure 4****Procedure 5**

5

In coupling procedure 1, an appropriate A domain (e.g., piperazine) is coupled to B domain (e.g., D-Boc-p-Cl-Phe-OH) in the presence of EDC/HOBt followed by Boc deprotection. The coupled AB compound is then coupled to an appropriate C domain followed by deprotection of Boc group and salt formation. Alternatively, when C domain is not protected with Boc group, the final compound can be obtained without the deprotection step.

In coupling procedure 2, an appropriate A domain (e.g., piperazine) is coupled to an appropriate BC domain in the presence of HATU followed by deprotection

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- 5 of Boc group and salt formation. Alternatively, when BC domain is not protected with Boc group, the final compound can be obtained without the deprotection step.

In coupling procedure 3, an appropriate AB domain is coupled to an appropriate C domain in the presence of EDC/HOBt followed by deprotection of Boc group and salt formation.

- 10 In coupling procedure 4, an appropriate BC domain is coupled to an appropriate A domain in the presence of EDC/HOBT followed by deprotection of Boc group and salt formation. Alternatively, when C domain is not protected with Boc group, the final compound can be obtained without the deprotection step.

- In coupling procedure 5, an appropriate AB domain is coupled to an
15 appropriate C domain in the presence of HATU followed by deprotection of Boc group salt formation.

For coupling of A with Boc-B, EDC/HOAT, EDC/HOBT or DCC/HOBT can be used.

- Generally, the starting material of Boc-protected piperazine (A domain)
20 can be deprotected in the presence of TFA/CH₂Cl₂, HCl/EtOAc, HCl/dioxane, or HCl in MeOH/Et₂O with or without a cation scavenger, such as dimethyl sulfide (DMS) before being subjected to the coupling procedure. It can be freebased before being subjected to the coupling procedure or in some cases used as the salt.

- A suitable solvent such as CH₂Cl₂, DMF, THF or a mixture of the above
25 solvents can be used for the coupling procedure. Suitable base includes triethyl amine (TEA), diisopropylethyl amine (DIPEA), N-methylmorpholine, collidine, or 2,6-lutidine. Base may not be needed when EDC/HOBT is used.

- Generally after the reaction is completed, the reaction mixture can be diluted with an appropriate organic solvent, such as EtOAc, CH₂Cl₂, or Et₂O, which is
30 then washed with aqueous solutions, such as water, HCl, NaHSO₄, bicarbonate, NaH₂PO₄, phosphate buffer (pH 7), brine or any combination thereof. The reaction mixture can be concentrated and then be partitioned between an appropriate organic solvent and an aqueous solution. The reaction mixture can be concentrated and subjected to chromatography without aqueous workup.

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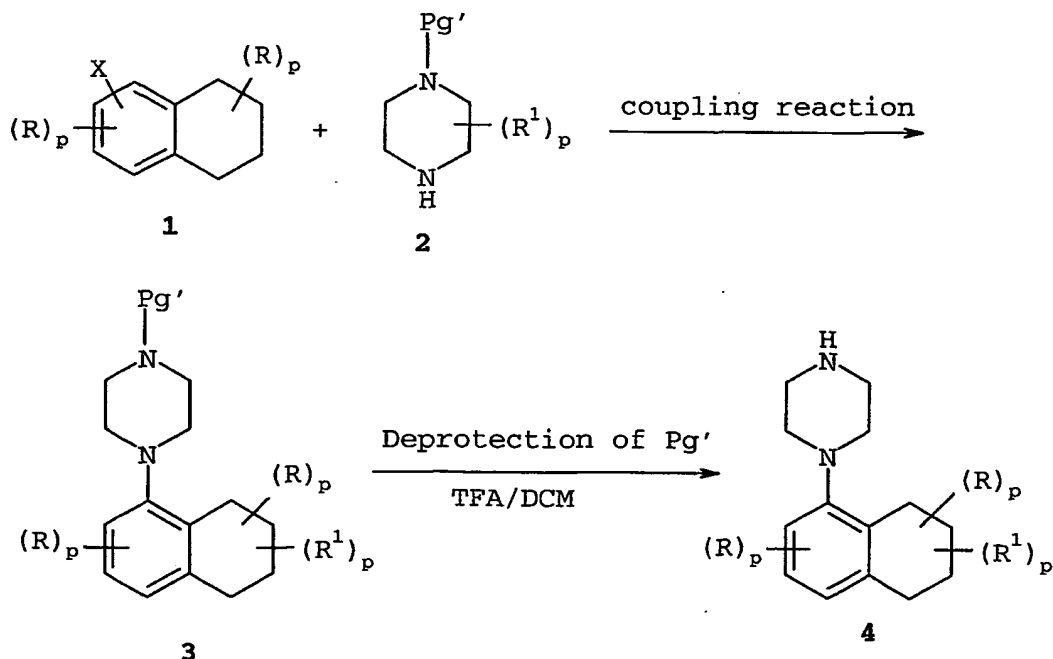
- 5 Protecting group such as Boc or CBZ, Fmoc, CF_3CO and $\text{H}_2/\text{Pd-C}$ can be deprotected in the presence of $\text{TFA}/\text{CH}_2\text{Cl}_2$, HCl/EtOAc , $\text{HCl}/\text{dioxane}$, HCl in $\text{MeOH}/\text{Et}_2\text{O}$, NH_3/MeOH , or TBAF with or without a cation scavenger, such as thioanisole, ethane thiol and dimethyl sulfide (DMS). The deprotected amines can be used as the resulting salt or are freebased by dissolving in CH_2Cl_2 and washing with
- 10 aqueous bicarbonate or aqueous NaOH . The deprotected amines can also be freebased by ion exchange chromatography.

The compounds of the present invention can be prepared as salt, such as TFA, hydrochloride or succinate salts by using known standard methods.

Preparation of "A domain"

- 15 The A domains of the present invention, in general, may be prepared from commercially available starting materials via known chemical transformations. The synthesis of certain A domains of the present invention is illustrated in Reaction Scheme 2 below.

20 Reaction Scheme 2: "A Domain"



X is halo; and

Pg' is protecting group such as BOC, CBZ or Fmoc.

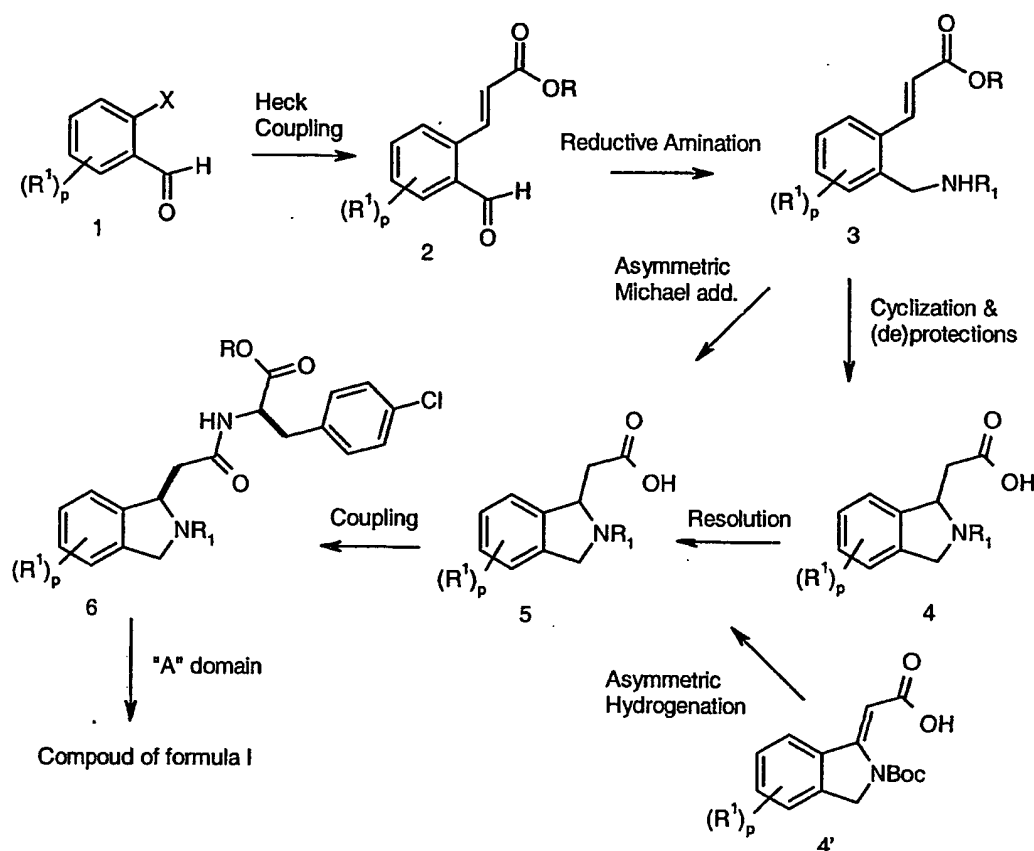
-75-

5 As shown in reaction scheme above, a compound of A domain (4) can be prepared by coupling benzofused carbocyclic compound (1) with piperazine (2) followed by deprotection of compound (3). A domain containing benzofused heterocyclyl can also be prepared in accordance with the reaction scheme 2.

10 The protected amino acid derivatives corresponding to the B and C domains are, in many cases, commercially available. Other protected amino acid derivatives can be prepared by known literature methods (see Williams, R. M. *Synthesis of Optically Active alpha-Amino Acids*, Pergamon Press: Oxford, 1989). More detailed examples of A Domain preparation are described below.

5 The present invention also provides a novel process for preparing certain intermediates and/or compounds of the invention as shown in Reaction Schemes 3-5.

Reaction Scheme 3:



10 As shown in Reaction Scheme 3, a convergent synthesis of a key intermediate isoindoline (5) via a Heck coupling, followed by a reductive amination, a ring cyclization and a resolution has been developed. Also, alternate asymmetric approaches including asymmetric Michael addition and asymmetric hydrogenation have also been developed to prepare compounds of the invention and/or intermediates thereof.

15 As shown in Reaction Scheme 3, the isoindoline compounds of the present invention may be prepared from 2-halobenzaldehyde 1 or substituted analog thereof. Preferred starting material is 2-bromobenzaldehyde or substituted analog thereof. Pd-mediated Heck coupling of 2-bromobenzaldehydes 1 with for example, methyl acrylate, provided α,β -unsaturated methyl esters 2, which undergoes reductive amination to

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5 give amines, 3 (or carbamates where R₁ is for example, Boc). Various Heck coupling reagents and conditions were found suitable to effect the coupling reaction. Suitable catalysts and ligands include Pd(OAc)₂/PPh₃, Pd(OAc)PPh₃/BU₄NBr, Pd(PPh₃)₂Cl₂/CUI, Pd(OAc)₂/P(O-Tol)₃. Suitable solvent or solvent systems for the Heck coupling reaction include DMF, toluene and ethyl acetate. More preferred base is triethylamine.

10 Reductive amination of the aldehyde functionality of 2 to amines is accomplished in good yields by reaction with benzylamine or alpha-methylbenzylamine in acidic conditions, followed by *in situ* reduction of the incipient imines with NaCNBH₃ at about pH 5. Other reducing agents including Na(OAc)₃BH and NaBH₄/H may also be used to effect reduction of the incipient imines. Interestingly, the resulting amines
15 immediately cyclized to the isoindoline compounds under the same acidic conditions conditions for the reduction. Direct preparation of compound 4 may also be effected by use of BocNH₂ instead of benzylamine in the reductive amination step. Screening of various reducing agents demonstrated that the combination of Et₃SiH and TFA in CH₃CN represents the preferred method for effecting reductive amination using BocNH₂.

20 The *N*-Boc isoindolinecarboxylic acid 5 may also be prepared from 3 as the carbamate, by an intra-molecular Michael addition and ester hydrolysis. The resolution of the isoindolinecarboxylic acids 4 by crystallization afforded enantio-pure compounds 5.

Two alternate asymmetric approaches have also been developed for the
25 synthesis of isoindolinecarboxylic acid 5 i.e., asymmetric Michael additions and asymmetric hydrogenation. In the asymmetric Michael addition approach, alpha-methylbenzyl amine is used as a chiral auxiliary to induce the enantio-selectivity. In the asymmetric hydrogenation approach, compound 4' could be converted to 5 stereoselectively in the presence of chiral ligands.

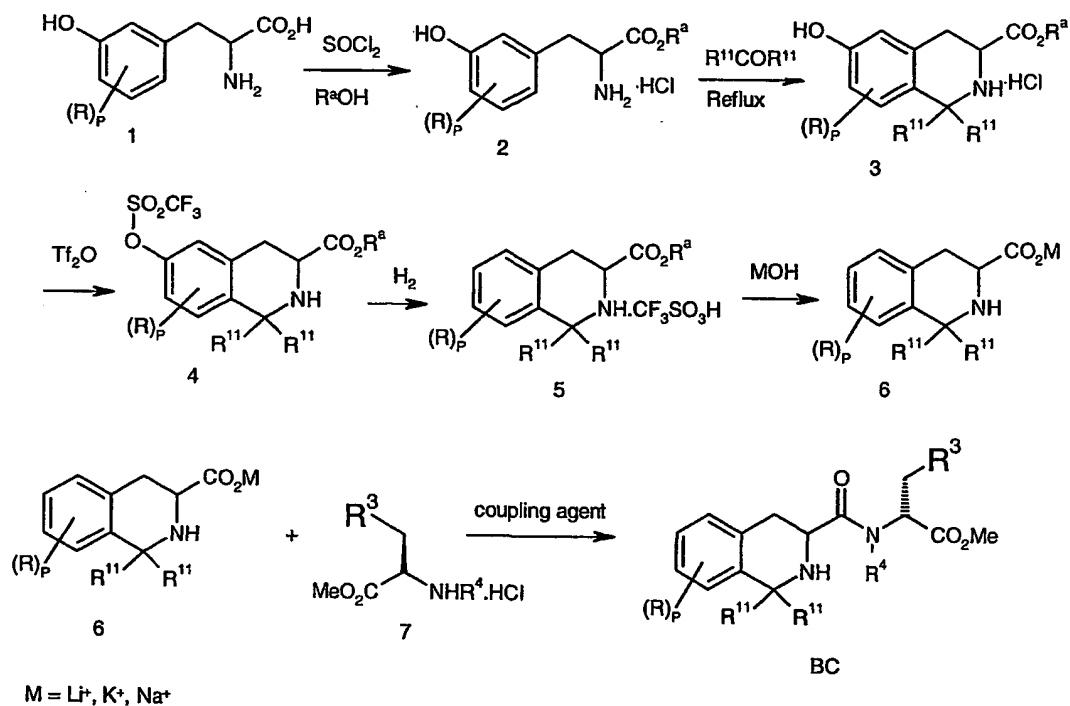
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5 Finally the coupling of the isoindolines 5 with the "B" domain piece, i.e., D-Cl-Phe to afford compound 6 ("BC" piece) is accomplished by standard amino acid coupling reactions such as, for example, by the use of EDC or EDCI or other activating agents in the presence of suitable dimethylaminopyridine (DMAP). The product (6) is then coupled with an "A" domain piece to afford the target MC4R agonist compound of formula I by coupling reactions known to one of skill in the art.

10 Preferably, the isoindole or other "C" domain piece is coupled to an "AB" coupled domain piece to form the compound of formula I.

Reaction Scheme 4



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5 As shown in Reaction Scheme 4, m-tyrosine ester or analogs, including substituted analogs thereof, may be esterified by forming the acid halide followed by nucleophilic displacement of halide by the alkoxy group from an alcohol, i.e., methanol or ethanol. Where thionyl chloride or other halide source is used the product may be isolated as the acid addition salt (2). The resulting ester (2) is subjected to a Pictet-Spengler reaction by heating with a suitable ketone or aldehyde in refluxing conditions. For example, an unsubstituted isoquinoline backbone (3) may be formed by employing formaldehyde in the Pictet-Spengler reaction. On the other hand, a gem-dimethyl substituted isoquinoline wherein R¹¹ is methyl, may be formed by using acetone as the ketone source and solvent. Other less reactive substituents may be substituted as the R¹¹ group for the practice of the present invention.

 The product isoquinoline (3) may be isolated preferably as the acid addition salt. Where m-tyrosine is used as the starting material, the free hydroxyl group is removed first by protection/activation with a good leaving group such as, for example, reaction with triflic anhydride (trifluoromethane sulfonic anhydride) or methanesulfonic acid to form the triflate or mesylate in the presence of a base. The triflate is a preferred group used to set up the compound (3) for deoxygenation because of the extra electron withdrawing effect of the trifluoromethane substituent. The deoxygenation reaction is effected by hydrogenation at pressures of about 50psi. The product (4) may be isolated as the acid addition salt. The product (4) is hydrolyzed under basic conditions to afford the acid salt. Suitable bases for the above hydrolysis include aqueous sodium hydroxide, potassium hydroxide and sodium lithium hydroxide. The reaction is preferably performed in a mixture of aqueous and organic solvents. An exotherm during addition of base may be regulated (i.e., less than about 35°C) to avoid overheating or "runaway reactions." The reaction product may be isolated by aqueous work up. Alternatively, the entire mixture may be concentrated and washed with organic solvents to afford the desired product (6) after crystallization.

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5 The product (6) is then reacted with a "B" domain substrate such as, for example, 4-chloro-D-phenylalanine as described previously and in the experimental section. The resulting "BC" combination product is then reacted with an "A" domain piece to form the respective compound of formula I. Alternatively, the product (6) may be reacted with an "AB" domain combination product to afford a compound of formula I.

10 One of skill is aware that certain protections and deprotections of intermediates in Reaction Scheme 4, to form the carbamate, substituted amine or free amine at the isoquinolinylnitrogen are possible and contemplated as within the scope of this invention. Unless otherwise specified, reagents and procedures for effecting the reactions described herein are known to one of skill in the art and may be found in
15 general reference texts such as *Advanced Organic Chemistry* by J. March, 5th edition, Wiley Interscience Publishers, New York, NY, and references therein.

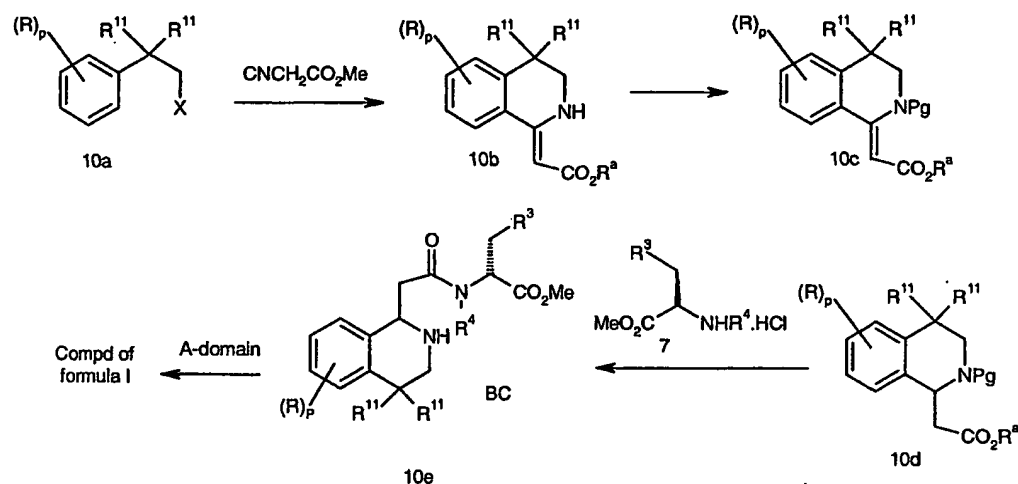
 In an alternate procedure, the isoquinoline product i.e., compound (3) or (5) including their N-protected analogs may be resolved by reaction with a resolving agent such as for example, L-tartaric acid, dehydroabietylamine or other resolving agents
20 known to one of skill in the art.

 Alternatively, asymmetric analogs of product (6) may be prepared by using asymmetric starting materials. For example, L-DOPA may be used in place of m-tyrosine ester in reactions essentially similar to those described and illustrated in Reaction Scheme 4, and in the examples, to afford the asymmetric analog of compound (6).
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5 Tetrahydroisoquinoline acetic acid derivatives may be prepared and utilized as shown in Reaction Scheme 5 below:

Reaction Scheme 5



10

As shown in Reaction Scheme 5, a compound of formula 10a wherein X is halogen, preferably bromo or chloro, and R and R¹¹ are as defined previously, and which is obtained commercially or prepared from commercial starting materials is reacted with cyanomethylethylacetate to afford a compound of formula 10b. The compound of formula 10b may be protected as the compound 10c with a suitable protecting group (Pg) and then subjected to hydrogenation conditions including for example asymmetric hydrogenation to form a compound of formula 10d, which may be chiral (depending on hydrogenation conditions, i.e., asymmetric versus non-assymmetric hydrogenation). The compound of formula 10d or stereoisomer thereof, is reacted with a B-domain piece such as, for example, 4-chloro-D-phe to afford a BC piece (10e). The compound of formula 10e is then reacted with an A-domain piece to afford a compound of formula I. The details of the specific reaction steps are similar to or analogous to reactions taught herein, and in the experimental section. Furthermore, one of skill in the art is aware of that such intermediate reactions as hydrolysis and deprotection may be necessary to achieve optimum yields in certain steps of the scheme as shown. One of skill in the art is also

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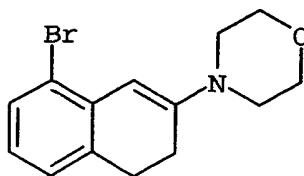
-82-

- 5 aware of further common manipulations such as N-alkylation, or N-acylation, and alkylations on the benzene ring to afford other compounds of formula I.

The following describes the detailed examples of A Domain preparation.

Preparation 1A

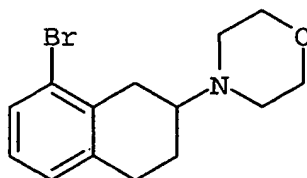
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- 15 To 8-Bromo-2-tetralone (3.0g, 13.3mmol) is added p-toluenesulfonic acid (0.25g, 1.47mmol), morpholine (7.43g, 85.3mmol), and toluene (25mL). The mixture is heated at reflux, using a Dean-Stark trap, for about 12 hours. After cooling to r.t. the mixture is concentrated to dryness and the crude enamine (3.9g) is taken on to the next step with no additional purification.

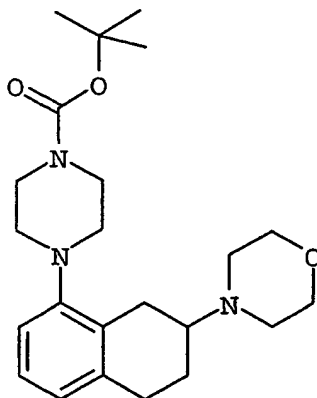
-83-

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Preparation 2A

To the compound of Preparation 1A (3.9g, 13.33mmol), in methanol (50mL), is added acetic acid (7.6mL, 133.3mmol) followed by careful addition of sodium borohydride (1.0g, 26.6mmol). The solution is allowed to stir at r.t. for about two hours. After concentrating to dryness, the remaining residue is taken up in EtOAc (100mL). The organics are washed with saturated NaHCO_3 (100mL), and the desired product is extracted into 1N HCl (100mL). The aqueous layer is then made alkaline with 5N NaOH (30mL) and extracted into EtOAc (100mL). The organic phase is concentrated to dryness to afford about 1.8g of the compound 2A (46%). EIS-MS 298.0 [M+1]

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Preparation 3A

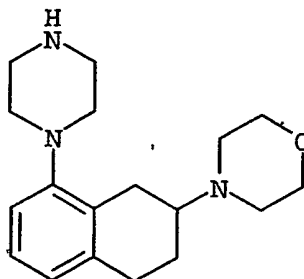
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To compound 2A (0.50g, 1.69mmol), in toluene (10mL), is added piperazine-1-carboxylic acid tert-butyl ester (0.314g, 1.69mmol), BINAP (0.168g, 0.27mmol), Na-*t*-BuO (0.243g, 2.53mmol) and $\text{Pd}_2(\text{dba})_3$ (0.124g, 0.135mmol). The solution is heated at reflux for about 4 hours, diluted with EtOAc (100mL), and filtered over a pad of celite. The filtrate is concentrated to dryness. The crude material is

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- 5 purified by flash chromatography (SiO₂, eluting with EtOAc to EtOAc-MeOH-TEA, 80:10:10) to afford about 0.65 g of the compound 3A (96%). EIS-MS 402.1 [M+1]

Preparation 4A



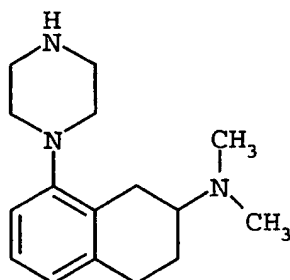
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To compound 3A (0.65g, 1.62mmol), in DCM (5mL), is added TFA (5mL) and the mixture is stirred at r.t. for about 45 minutes. After concentrating to dryness, the resulting residue is taken up with 1N NaOH and the desired "free" amine is extracted into EtOAc (100mL). The organic extracts are concentrated to dryness to afford

15 about 290mg of the compound 4A (60%). EIS-MS 302.0 [M+1]

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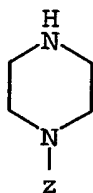
Preparation 5A

The 2-dimethylamino-8-bromo-1,2,3,4-tetrahydronaphthalene (1.0g, 3.93mmol) is prepared by following a substantially similar procedure as described in Preparation 1A and 2A, and from commercially available 1-Boc-piperazine (0.805g, 4.32mmol) as described *J. Am. Chem. Soc.*, 118:7215-7216, 1996. The desired product is purified by flash chromatography. Yield: 1.35g (95%). EIS MS: 360.2 [M+H].

To the Boc-protected compound (1.35g, 3.75mmol) is added TFA (5mL) and DCM (5mL) and the resulting mixture is stirred at r.t. for about 30 minutes. After concentrating to dryness, the resulting residue is taken up in 1N NaOH and the desired amine extracted into EtOAc (100mL). The organics are washed with H₂O and brine, and then re-concentrated. Yield: 970 mg. EIS-MS: 260.1 [M+H]

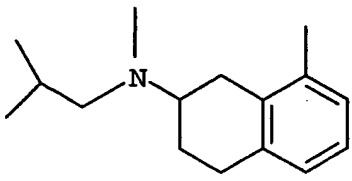
-86-

- 5 The A domain compounds 6A to 11A are prepared from 8-bromo-2-tetralone and an appropriate amine by following the procedure substantially similar to that described in Preparation 4A.

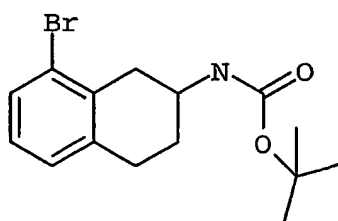


No	Z	EIS-MS [M+1]
6A	<p>Chemical structure of 1-(1,2,3,4-tetrahydronaphthalen-1-yl)diethylamine, where the nitrogen atom is substituted with two ethyl groups.</p>	385.3
7A	<p>Chemical structure of 1-(1,2,3,4-tetrahydronaphthalen-1-yl)diethylamine, where the nitrogen atom is substituted with two propyl groups.</p>	—
8A	<p>Chemical structure of 1-(1,2,3,4-tetrahydronaphthalen-1-yl)pyrrolidine, where the nitrogen atom is part of a pyrrolidine ring.</p>	286.2
9A	<p>Chemical structure of 1-(1,2,3,4-tetrahydronaphthalen-1-yl)N-ethyl-N-methylamine, where the nitrogen atom is substituted with an ethyl group and a methyl group.</p>	274.2
10A	<p>Chemical structure of 1-(1,2,3,4-tetrahydronaphthalen-1-yl)N-isopropyl-N-methylamine, where the nitrogen atom is substituted with an isopropyl group and a methyl group.</p>	288.2

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No	Z	EIS-MS [M+1]
11A		302.2

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Preparation 12A

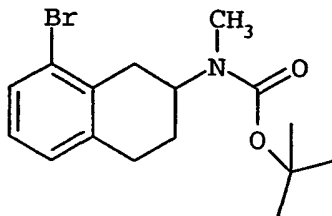
To commercially available 8-bromo-2-aminotetralin (0.85g, 3.76mmol) is added Boc_2O (0.90g, 4.13mmol), K_2CO_3 (1.03g, 7.52mmol) and THF/ H_2O (1:1, 25mL).

- 10 The mixture is stirred for about 30 minutes and then diluted 10 fold with EtOAc. The organics are washed with saturated NaHCO_3 , H_2O , and brine, and then concentrated to dryness. Material is recrystallized from hexanes to afford about 1.2g of compound 12A (98%). EIS-MS 327.1 [M+1]

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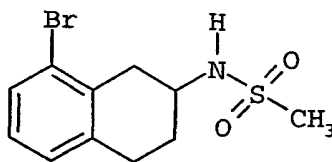
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Preparation 13A

To compound 12A (0.18g, 0.54mmol) in THF (3mL) is added NaH (60% dispersion in oil, 0.03g, 0.80mmol). The solution is allowed to stir at r.t. for about one
10 hour. Iodomethane (0.15g, 1.07mmol) is then added drop-wise to the solution. The mixture is allowed to stir at r.t. for about 18 hours. This mixture is then diluted 10 fold with EtOAc (30mL) and washed with water (30mL). The organic phase is concentrated to dryness to afford about 0.13g of compound 13A (69%). EIS-MS 340.1 [M+1]

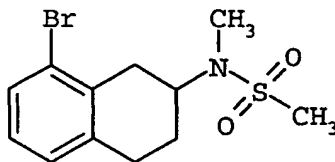
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Preparation 14A

To commercially available 8-bromo-2-aminotetralin (0.26g, 1.14mmol) in
CH₂Cl₂ (5mL), is added pyridine (0.28mL, 3.44mmol), and methanesulfonyl chloride
20 (0.089mL, 1.14mmol). The mixture is stirred at r.t. for about 30 minutes and then diluted with EtOAc (50mL). The organics are washed with H₂O and brine and then concentrated to dryness to afford about 180 mg of compound 14A (52%). EIS-MS 304.9 [M-1]

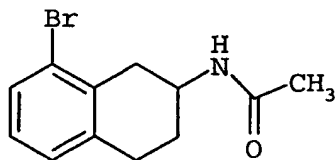
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Preparation 15A

To compound 14A (0.275g, 0.903mmol), in THF (5mL) is added NaH (60% dispersion in oil, 0.038g, 1.54mmol). The solution is allowed to stir at r.t. for about one hour. Iodomethane (0.061mL, 0.99mmol) is added drop-wise to the solution. The reaction mixture is allowed to stir at r.t. for about 12 hours. This mixture is then diluted 10 fold with EtOAc (30mL) and washed with water (30mL). The organic phase is concentrated to dryness. The desired product is purified by flash chromatography (SiO₂, eluting with 30% EtOAc in Hexanes) to afford about 0.227g of compound 15A (79%).

15 EIS-MS 320.1 [M+1]

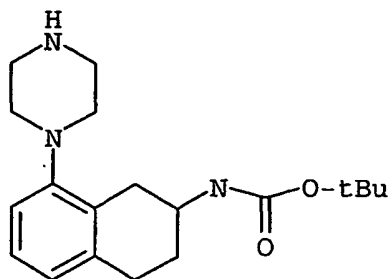
Preparation 16A

To commercially available 8-bromo-2-aminotetralin (1.5g, 6.63mmol) is added Ac₂O (10mL) and pyridine (10mL). The mixture is heated at 80°C for about 4 hours. After cooling to r.t., the mixture is concentrated to dryness and the resulting residue taken up in EtOAc (50mL). The organics are washed with saturated NaHCO₃, H₂O and brine, and then concentrated to dryness. The product is recrystallized from EtOAc-Hexanes to afford about 1.75g of compound 16A (98%). EIS-MS 270.1 [M+1]

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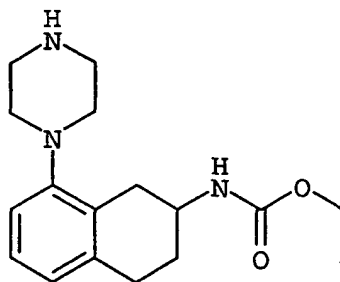
Preparation 17A

To commercially available 8-bromo-2-aminotetralin (850 mg, 3.76mmol) is added Boc_2O (900 mg, 4.13mmol), K_2CO_3 (1.03g, 7.52mmol) and THF/ H_2O (1:1, 25mL). The mixture is rapidly stirred for about 30 minutes and then diluted 10 fold with EtOAc. The organics are washed with saturated NaHCO_3 , H_2O and brine, and then concentrated to dryness. The material is recrystallized from hexanes. Yield: 1.2 g (98%).

To the compound obtained above (300mg, 0.920 mmol), piperazine is added (316 mg, 3.68 mmol) by following conditions outlined in *J. Am Chem. Soc.*, 118:7215-7216 for the coupling. The final compound is purified by flash chromatography (SiO_2 by eluting with 90:5:5, EtOAc- Et_3N -MeOH) to afford about 248g of compound 17A (81%). EIS-MS: 332.2 $[\text{M}+1]$

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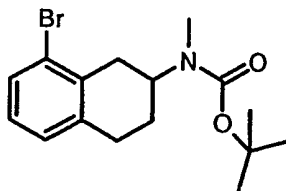
Preparation 18A

To 2-amino-8-(4-t-butyloxycarbonylpiperazin-1-yl)decalin (75mg, 0.226mmol), in DCM (5mL) is added Et₃N (63 microliters) and methyl chloroformate (17 microliters). The mixture is stirred at r.t. for about 30 minutes and then concentrated to dryness. The resulting residue is taken up in EtOAc (50mL) and the organics are washed with saturated NaHCO₃, H₂O and brine, and then concentrated to dryness. Desired product is purified by flash chromatography (SiO₂ by eluting EtOAc. Yield: 40 mg (45%). EIS-MS: 390.2 [M+H]

Following the procedure as described in Preparation 5A, the Boc-protected compound from above (40mg, 0.103mmol) is deprotected to afford about 26 mg (87%) of compound 18A. EIS-MS: 290.1 [M+H]

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Preparation 19A

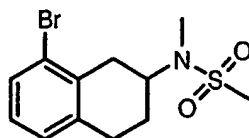
Step 1: To commercially available 8-bromo-2-aminotetralin (0.85 g, 3.76 mmol) is added Boc_2O (0.90 g, 4.13 mmol), K_2CO_3 (1.03g, 7.52 mmol) and THF/ H_2O (1:1, 25 mL). The mixture is stirred for about 30 minutes and then diluted 10 fold with EtOAc. The organics are washed with saturated NaHCO_3 , H_2O and brine, and then concentrated to dryness. Material is recrystallized from hexanes yielding an off white solid (1.2 g, 98%). EIS-MS 327.1 $[\text{M}+1]$

Step 2: To the compound from Step 1 (0.18 g, 0.54 mmol) in THF (3 mL) is added NaH (60% dispersion in oil, 0.03 g, 0.80 mmol). The solution is allowed to stir at r.t. for about an hour. Iodomethane (0.15 g, 1.07 mmol) is then added dropwise to the solution. The reaction mixture is allowed to stir at r.t. for about 18 hours. This mixture is then diluted 10 fold with EtOAc (30 mL) and washed with water (30 mL). The organic phase is then concentrated to dryness to afford about 0.13 g of the final compound (69%). EIS-MS 340.1 $[\text{M}+1]$

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Preparation 20A

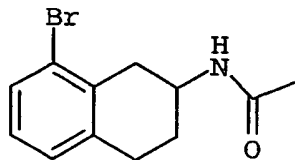
Step 1: To commercially available 8-bromo-2-aminotetralin (0.26 g, 1.14 mmol) in DCM (5 mL) is added pyridine (0.28 mL, 3.44 mmol), and methanesulfonyl chloride (0.089 mL, 1.14 mmol). The mixture is stirred at r.t. for about 30 minutes and then diluted with EtOAc (50 mL). The organics are washed with H₂O and brine and then concentrated to dryness. Yield: 180 mg, 52%. EIS-MS 304.9 [M-1]

Step 2: To compound from step 1 (0.275 g, 0.903 mmol), in THF (5 mL) is added NaH (60% dispersion in oil, 0.038 g, 1.54 mmol). The solution is stirred at r.t. for about an hour. Iodomethane (0.061 mL, 0.99 mmol) was then added dropwise to the solution. The mixture is stirred at r.t. for 12 hours. This mixture is then diluted 10 fold with EtOAc (30 mL) and washed with water (30 mL). The organic phase is concentrated to dryness. The desired product is purified by flash chromatography (SiO₂, eluting with 30% EtOAc in Hexanes) to afford about 0.227 g of the final compound (79%). EIS-MS 320.1 [M+1]

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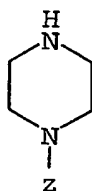
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Preparation 21A

To commercially available 8-bromo-2-aminotetralin (1.5 g, 6.63 mmol) is added Ac₂O (10 mL) and pyridine (10 mL). The mixture is heated at 80°C for about 4 hours. After cooling to r.t., the mixture is concentrated to dryness and the resulting residue was taken up in EtOAc (50 mL). The organics is washed with saturated NaHCO₃, H₂O and brine and then concentrated to dryness. The product is recrystallized from EtOAc-hexanes to afford about 1.75g of the final compound (98%). EIS-MS 270.1 [M+1]

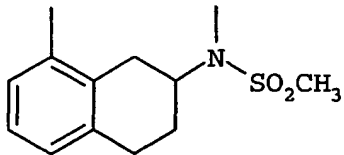
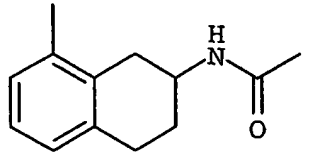
Preparation of 19Aa-21Aa

The A domain compounds of 19Aa-21Aa are prepared from the appropriately substituted 8-bromo-2-tetraline and N-Boc-piperazine by following substantially similar procedures as described in Preparations 3 to 4.

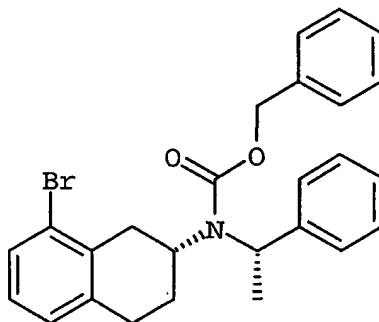


No	Z	EIS-MS [M+1]
19Aa		346.2

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20Aa		324.1
21Aa		274.1

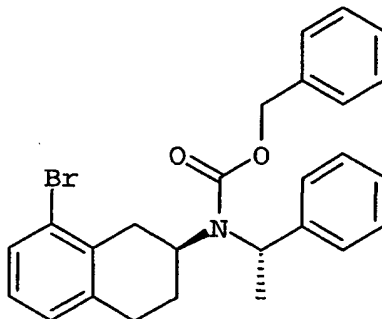
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Preparation 22A

To (15.31 g, 46.35 mmol) of 8-bromo-1,2,3,4-tetrahydro-naphthalen-2-yl)-(1-phenyl-ethyl)-amine, whose preparation was described in *J. Med. Chem.* 1993, 36(15), 2066-2074, is added THF (50 mL), H₂O (50 mL), K₂CO₃ (12.81g, 92.71 mmol) and the mixture is cooled to 0°C. Benzyl chloroformate (6.61mL, 46.35 mmol) is then added and the mixture is allowed to warm to r.t. and stirred for an additional 30 minutes. After diluting with EtOAc (5 fold), the organics are washed with H₂O, saturated NaHCO₃, brine, and concentrated to dryness. The resulting residue is purified by flash chromatography (SiO₂, eluting with 10%EtOAc in Hexanes) to afford about 17.5g of the final compound (81%). EIS-MS 464.2 [M+1]

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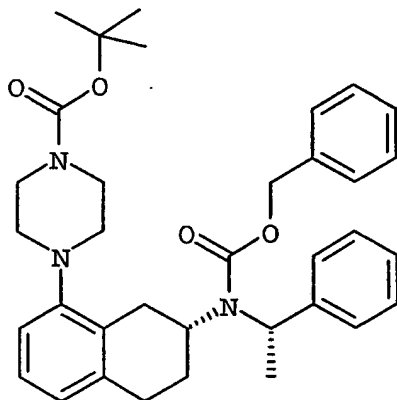
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Preparation 23A

To (1.54, 4.66 mmol) of 8-bromo-1,2,3,4-tetrahydro-naphthalen-2-yl)-(1-phenyl-ethyl)-amine, whose preparation is described in *J. Med. Chem.* **1993**, 36(15), 2066-2074, is added THF (10 mL), H₂O (10 mL), K₂CO₃ (1.28, 92.71 mmol) and the mixture is cooled to 0°C. Benzyl chloroformate (0.79 mL, 5.59 mmol) is then added and the mixture is allowed to warm to r.t. and stirred for an additional 30 minutes. After diluting with EtOAc (5 fold), the organics are washed with H₂O, saturated NaHCO₃, brine, and concentrated to dryness. The resulting residue is purified by flash chromatography (SiO₂, eluting with 10%EtOAc in Hexanes) to afford about 2.0g of the final compound (92%). EIS-MS 464.2 [M+1]

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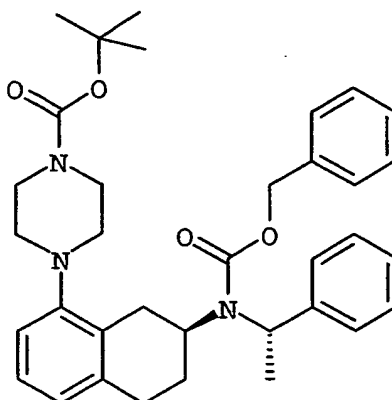
Preparation 24A

The compound 24A is prepared from the compound 22A (17.5 g, 37.7 mmol) by essentially following the procedure as described in Preparation 3A to afford about 15.86g (74%). EIS-MS 570.2 [M+1]

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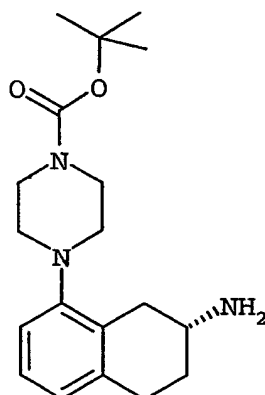
-98-

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Preparation 25A

The compound 25A is prepared by essentially following the procedure as described in Preparation 23A to afford about 1.28g (52%). EIS-MS 570.2 [M+1]

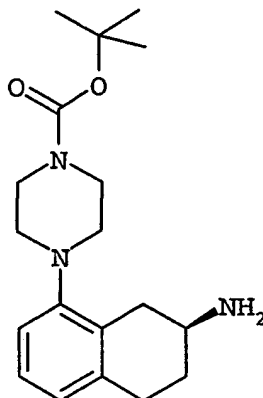
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Preparation 26A

To Pd(OH)₂/C (7.5g), 10%Pd/C (15g), in EtOH/HOAc (4:1, 125 mL), is added the compound 24A (15.85 g, 27.8 mmol) in EtOH/HOAc (4:1, 125 mL), and the mixture is stirred at r.t. under 1 atm H₂ overnight. The catalyst is removed by filtering over a bed of celite and the filtrate is concentrated to dryness. The resulting residue is purified by flash chromatography (SiO₂, eluting with 90:5:5, EtOAc-TEA-MeOH) to afford about 6.5g of the final compound (71%). EIS-MS 332.3 [M+1]

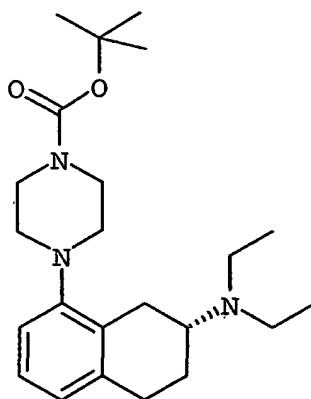
-99-

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Preparation 27A

To $\text{Pd}(\text{OH})_2/\text{C}$ (0.6g), 10%Pd/C (1.28g), in EtOH/HOAc (4:1, 40 mL), is added the compound 25A (1.28 g, 2.25 mmol) in EtOH/HOAc (4:1, 30 mL), and the mixture is stirred at r.t. under 1 atm H_2 overnight. The catalyst is removed by filtering over a bed of celite and the filtrate is concentrated to dryness. The resulting residue is purified by flash chromatography (SiO_2 , eluting with 90:5:5, EtOAc-TEA-MeOH) to afford about 340 mg of the final compound (46%). EIS-MS 332.3 [M+1]

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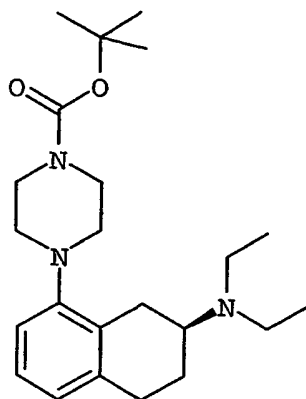
Preparation 28A

To the compound 26A (2.0 g, 6.03 mmol) is added K_2CO_3 (4.16 g, 30.16 mmol), ethyl bromide (1.35 mL, 18.09 mmol) and DMF (1 mL). The mixture is stirred at r.t. for about 24 hours and then diluted 10 fold with EtOAc. The organics are washed with H_2O , brine and concentrated to dryness. The resulting residue is purified by flash

-100-

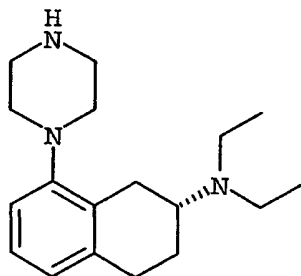
- 5 chromatography (SiO_2 , eluting with 98:1:1, EtOAc-TEA-MeOH) to afford about 1.86g of the final compound (80%). EIS-MS 388.4 [M+1]

Preparation 29A



- 10 Compound 29A is prepared from the compound 27A by following the procedure as described in Preparation 28A. EIS-MS 388.4 [M+1]

Preparation 30A

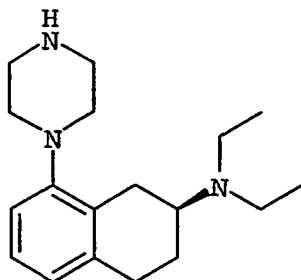


- 15 To the compound 28A (0.35g, 0.903 mmol) in DCM (5 mL) is added TFA (5 mL), and the mixture is stirred at r.t. for about 45 minutes. After concentrating to dryness, the resulting residue is taken up with 1N NaOH and the desired "free" amine is extracted into EtOAc (100 mL). The organic extracts are concentrated to dryness to afford about 230mg of the final compound (89%). EIS-MS 288.2 [M+1]

20

Preparation 31A

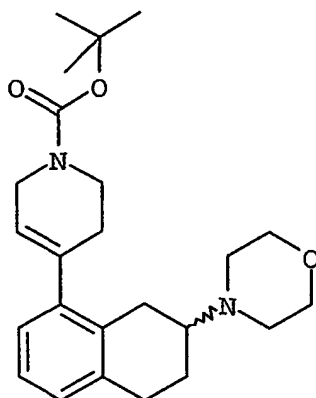
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Compound 31A is prepared from the compound 29A by following a substantially similar procedure as described in Preparation 30A. EIS-MS 288.2 [M+1]

Preparation 32A



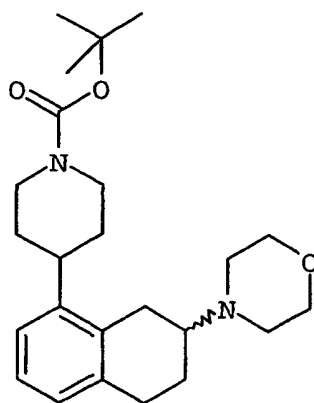
10

To the compound 2A (0.725 g, 2.45 mmol), in DMF (20 mL), is added 1-2(H)-pyridine-carboxylic acid, 3,6-dihydro-4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1,1-dimethyl ethyl ester (0.909 g, 2.94 mmol, *Tet. Lett.*, 41:3705-3708, 2000), dichloro(1,1'-bis(diphenylphosphino)ferrocene)palladium (II) DCM adduct (0.108 g, 0.147 mmol), and K_2CO_3 (1.02 g, 7.35 mmol). The reaction is heated to about 90°C overnight. The mixture is cooled, diluted with DCM and filtered through celite. The filtrate is concentrated to dryness and the resulting residue is taken up in EtOAc (50 mL). The organics are washed with H_2O , brine, and concentrated to dryness. The desired product is purified by flash chromatography (SiO_2 , eluting with 100% EtOAc to EtOAc-TEA-MeOH, 80:10:10) to afford about 624mg of the final compound (64%). EIS-MS 399.2 [M+1]

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Preparation 33A

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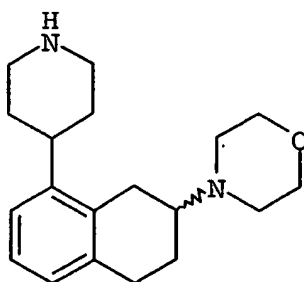
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To the compound 32A (0.62 g, 1.57 mmol) in EtOH (20 mL) is added a slurry of 10% Pd/C (0.62 g) in EtOH (20 mL). The mixture is stirred rapidly under H₂ (1 atm) for about 2 hours. The reaction mixture is filtered over a pad of celite and washed with EtOAc (100 mL). The filtrate concentrated to dryness to afford about 0.58g of the final compound (83%). EIS-MS 401.3 [M+1]

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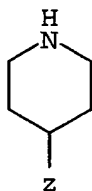
-103-

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Preparation 34A

To the compound 33A (0.50 g, 1.25 mmol) in DCM (3 mL) is added TFA (3 mL), and the mixture is stirred at r.t. for about 45 minutes. After concentrating to dryness, the resulting residue is taken up with 1N NaOH and the desired “free” amine
 10 extracted into EtOAc (50 mL). The organic extracts are concentrated to dryness to afford about 300 mg of the final compound (81%). EIS-MS 301.2 [M+1]

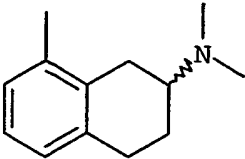
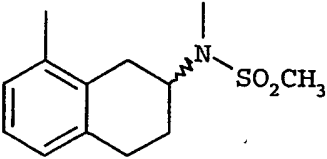
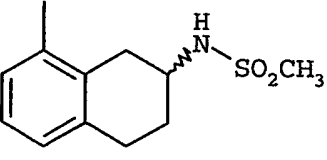
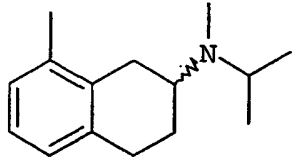
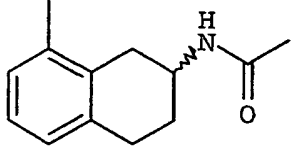
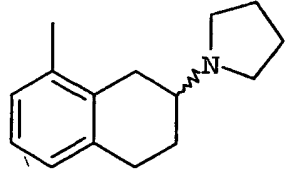
The A domain 35A–42A are prepared from using appropriate 8-bromo-2-tetralin (described in Preparation 1A and 2A) by following a substantially similar procedures as described in Preparations 32A–34A.



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No	Z	(EIS)-MS [M+1]
35A		287.2
36A		315.3

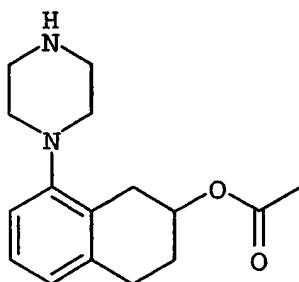
-104-

No	Z	(EIS)-MS [M+1]
37A		--
38A		323.1
39A		309.1
40A		287.2
41A		273.1
42A		285.2

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Preparation 43A



Step 1: 8-Bromo-2-hydroxytetralin is prepared from 8-bromo-2-tetralone (3.0 g, 13.3mmol) as described in *J. Org. Chem.*, 56(19):5564-5566, 1991. The desired product is purified by flash chromatography (Silica gel 60, eluting with 25% EtOAc in Hexanes) to afford about 2.7 g (90%). EIS-MS: 228 [M+H].

Step 2: To 8-bromo-2-hydroxytetralin (2.0 g, 8.81mmol) is added TBSCl (1.39 g, 9.25mmol), imidazole (1.19g, 17.6mmol) and DMF (20mL). The mixture is stirred at r.t. for about 5 hours and then diluted 10 fold with EtOAc. The organics are washed with H₂O and brine, and then concentrated to dryness to afford about 2.5 g (83%).

15 Step 3: 1-Boc-piperazine is coupled to the TBS-protected alcohol from above (2.49g, 7.29mmol) as described in *J.Org.Chem.*, 61:7240-7241, 1996. The desired product is purified by MPLC to afford about 1.6g (50%). EIS-MS: 447.3 [M+H].

Step 4: To the coupled product (810 mg, 1.81mmol), in THF (10mL), is added tetrabutylammonium fluoride (1.0M in THF, 3.6mL, 3.6mmol) and the reaction mixture stirred at r.t. for about 18 hours. The mixture is diluted 10 fold with EtOAc, and the organics are washed with saturated NaHCO₃, H₂O and brine, and then concentrated to dryness to give about 540 mg of 1-t-butyloxycarbonyl-4-(2-hydroxytetralin-8-yl)piperazine (90%). EIS-MS: 332.2 [M+H].

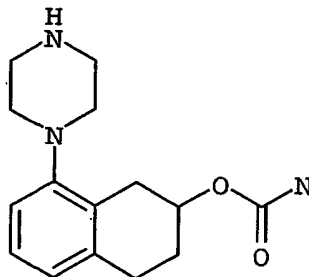
Step 5: To compound of Step 4 (96mg, 0.288mmol) is added Ac₂O (5mL) and pyridine (5mL). The mixture is stirred at 80°C for about 1.5 hours and then allowed to cool to r.t. After concentrating to dryness, the resulting residue is taken up in EtOAc (100mL) and the organics are washed with H₂O and brine, and then concentrated to dryness to give about 75 mg of Boc-protected compound (70%). EIS-MS: 375.2 [M+H].

Step 6: To the compound of step 5 (75 mg, 0.200mmol) is added TFA (4mL) and DCM (4mL). The resulting mixture is stirred at r.t. for about 1 hour and then

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- 5 concentrated to dryness. The resulting residue is taken up in H₂O at pH 10 (1N NaOH) and the desired amine is extracted into EtOAc (50mL). The organics are washed with H₂O and brine and then concentrated to dryness. EIS-MS: 275.2 [M+H].

Preparation 44A



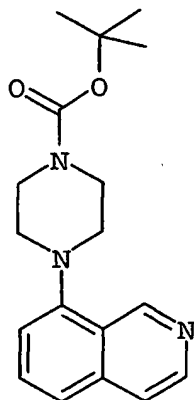
10

The Boc-protected compound is prepared from 1-t-butyloxycarbonyl-4-(2-hydroxytetralin-8-yl)piperazine (107mg, 0.321mmol) as described in *J.Chem.Perk.Trans.1*, 4:377-382, 1996 (85 mg, 71%). EIS-MS: 376.2 [M+H].

- 15 The final compound is prepared from the Boc-protected analogue (80mg, 0.213mmol) by following the procedure as described in Preparation 5A to afford about 58mg (100%) of the final compound. Ion spray MS: 276.1 [M+H].

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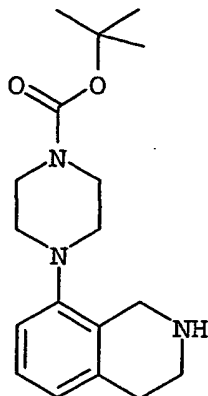
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Preparation 45A

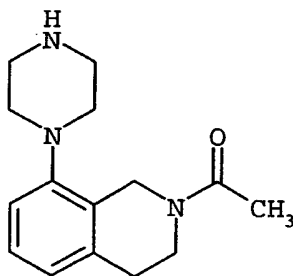
The 8-bromoisoquinoline (3.18 g, 15.28 mmol), piperazine (3.29 g, 38.21 mmol), Pd₂(dba)₃ (696 mg, 0.76 mmol), BINAP (1.43 g, 2.29 mmol) and sodium *tert*-butoxide (2.06 g, 21.39 mmol) are dissolved in toluene (80 mL), and the mixture is heated to 90°C for about 16 hours. The mixture is allowed to cool to r.t. and filtered across celite and concentrated. The residue is purified by silica gel chromatography, which is then dissolved in DCM (100mL). Excess BOC anhydride is added to the mixture while stirring. The mixture is allowed to react for about 2 hours and then concentrated. The product is purified by chromatography on silica gel to afford about 3.97g of the final compound (83 %). EIS-MS 214.2 [M-BOC].

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Preparation 46A

The compound 45A (1.2 g, 3.83 mmol) is dissolved in a slurry of PtO_2 (1 g) in isopropanol (150 mL), and pressurized to 45 psi with hydrogen gas overnight. The slurry is filtered and concentrated to dryness. The resultant oil is purified by flash chromatography on silica gel to afford about 920 mg of the final compound (76%). EIS-MS 318.1 [M+1]

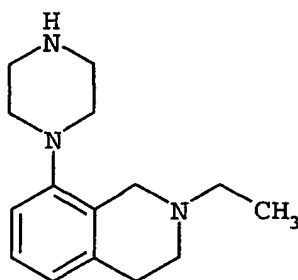
Preparation 47A

The compound 46A (150 mg, 0.47 mmol) is dissolved in acetic anhydride (4 mL) and TEA (1mL) is added. The mixture is stirred for about 4 hours. The mixture is then diluted with water and DCM (50 mL/50 mL) and the pH is adjusted to about 10 with 1M NaOH. The organic fraction is concentrated to dryness, and the resulting residue purified by flash chromatography on silica gel. The product is stirred in 1:1 TFA/DCM (10 mL) for about 2 hours. The mixture is concentrated to dryness and the resulting residue is subjected to SCX ion exchange chromatography followed by flash

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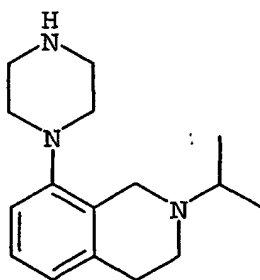
- 5 chromatography on silica gel to give about 113mg of the final product (93%). EIS-MS 260.1 [M+1]

Preparation 48A



- 10 The compound 47A (337 mg, 1.38 mmol) is dissolved in THF (10 mL) and lithium aluminum hydride (210 mg, 5.5 mmol) is added as a slurry in THF (5 mL). The mixture is stirred for about 4 hours at reflux. The reaction is worked up with water and the organics are washed with 1N NaOH and then concentrated to dryness. The resulting residue is purified by flash chromatography on silica gel to give about 226mg of
- 15 the final product (67%). EIS-MS 246.2 [M+1]

Preparation 49A

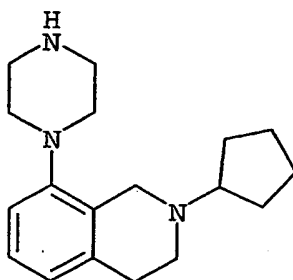


- To the compound 46A (250 mg, 0.79 mmol) in DCM (3.6 mL) is added
- 20 acetone (0.11 mL, 1.5mmol), sodium triacetoxyborohydride (469 mg, 2.21 mmol) and acetic acid (0.272 mL, 4.74 mmol), and the mixture is stirred at r.t. for about 24 hours. The mixture is diluted with DCM (50 mL) and washed with 1M NaOH. The organic fraction is concentrated and purified by silica gel chromatography, and the product stirred in 1:1 TFA/DCM (10 mL) for about 2 hours. The reaction is concentrated and the
- 25 resulting residue is subjected to SCX ion exchange chromatography followed by silica gel

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- 5 chromatography to give about 182mg of the final compound (89%). EIS-MS 260.1 [M+1]

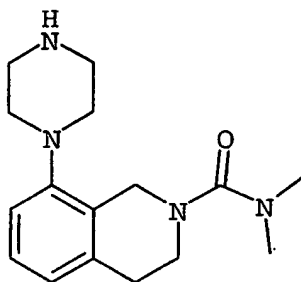
Preparation 50A



- 10 To the compound 46A (250 mg, 0.79 mmol) in DCM (3.6 mL) is added cyclopentanone (0.133 mL, 1.5 mmol), sodium triacetoxyborohydride (469 mg, 2.21 mmol) and acetic acid (0.272 mL, 4.74 mmol), and the reaction is stirred at r.t. for about 24 hours. The reaction is diluted with DCM (50 mL) and washed with 1M NaOH. The organic fraction is concentrated and purified by silica gel chromatography, and the
- 15 product stirred in 1:1 TFA/DCM (10 mL) for about 2 hours. The reaction is concentrated, and the resulting residue is subjected to SCX ion exchange chromatography followed by silica gel chromatography to give about 162 mg of the final product (79%). EIS-MS 286.2 [M+1]

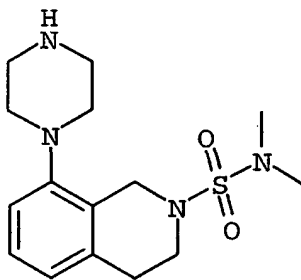
-111-

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Preparation 51A

To the compound 46A (400 mg, 1.26 mmol) in DCM (10 mL) is added dimethylcarbamyl chloride (0.329 mL, 3.78 mmol) and TEA (1.05 mL, 7.56 mmol), and the reaction is stirred at r.t. for about 16 hours. The mixture is diluted with DCM (50 mL) and washed with 1M NaOH. The organic fraction is concentrated and purified by silica gel chromatography, and the product is stirred in 1:1 TFA/DCM (10 mL) for about 2 hours. The reaction is concentrated, and the residue subjected to SCX ion exchange chromatography followed by silica gel chromatography to give about 371mg of the final product. EIS-MS 289.1 [M+1]

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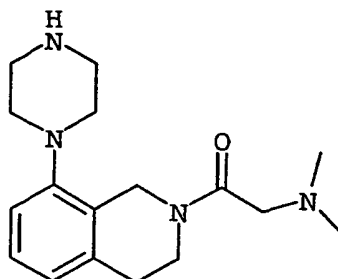
Preparation 52A

To the compound 46A (400 mg, 1.26 mmol) in DCM (10 mL) is added dimethylsulfamoyl chloride (0.406 mL, 3.78 mmol) and TEA (1.05 mL, 7.56 mmol), and the mixture is stirred at r.t. for about 16 hours. The mixture is diluted with DCM (50 mL) and washed with 1M NaOH. The organic fraction is concentrated and purified by silica gel chromatography, and the product stirred in 1:1 TFA/DCM (10 mL) for about 2 hours. The mixture is concentrated, and the residue subjected to SCX ion exchange chromatography followed by silica gel chromatography to give about 422mg of the final product. EIS-MS 325.1 [M+1]

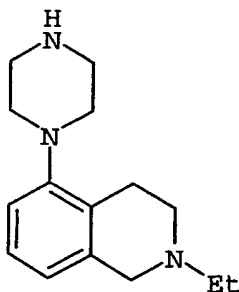
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Preparation 53A

To the compound 46A (400 mg, 1.26 mmol) in DCM (10 mL) is added *N,N*-dimethyl glycine (130 mg, 1.26 mmol), EDC (242 mg, 1.26 mmol), HOBT (170 mg, 1.26 mmol) and DIPEA (0.44 mL, 1.26 mmol), and the mixture is stirred at r.t. for about 16 hours. The mixture is diluted with DCM (50 mL) and washed with 1M NaOH. The organic fraction is concentrated and purified by silica gel chromatography, and the product is stirred in 1:1 TFA/DCM (10 mL) for about 2 hours. The reaction is concentrated, and the residue is subjected to SCX ion exchange chromatography followed by silica gel chromatography to give about 348 mg of the final product (91%). EIS-MS 303.2 [M+1]

Preparation 54A

Step 1: 5-Bromoisoquinoline (250mg, 1.2mmol) is dissolved in toluene (5mL). To the solution is added piperazine (258mg, 3.0mmol), Pd₂dba₃ (55mg, 0.06mmol), BINAP (112mg, 0.18mmol), and sodium t-butoxide (162mg, 1.68mmol). The mixture is degassed and then heated to 90°C for about 12 hours. The reaction is allowed to cool to r.t., diluted with diethyl ether and filtered through celite. The filtrate is

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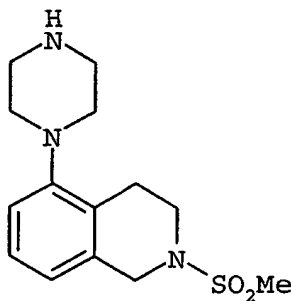
5 concentrated, and the dark residue is chromatographed on silica gel to give about 160 mg of 5-(piperazin-1-yl)isoquinoline (63%). EIS-MS: Found 214.0 (M+1).

Step 2: 5-(Piperazin-1-yl)isoquinoline is dissolved in DCM, and Boc₂O (620mg, 2.84mmol) is added in one portion. The reaction mixture is stirred at r.t. for about 4 hours under nitrogen and concentrated under vacuum. The residue is purified by
10 chromatography on silica gel to give about 201mg of 5-(4-t-butyloxycarbonylpiperazin-1-yl)isoquinoline (92%).

Step 3: The compound of Step 2 (200mg, 0.64mmol) is dissolved in absolute ethanol (50mL), and Pt/C 10% catalyst (200mg) is added. The mixture is pressurized to 45psi under hydrogen and shaken for about 4 hours at r.t. The mixture is
15 filtered over celite, concentrated and chromatographed on silica gel to give about 68mg of 5-(4-t-butyloxycarbonylpiperazin-1-yl)-2-ethyl-1,2,3,4-tetrahydroisoquinoline (30%).

Step 4: The compound of Step 3 (68mg, 0.196mmol) is dissolved in DCM (1.5mL) and TFA (1.5mL) and the mixture is stirred for about 1.5 hours. The mixture is concentrated to an oil, which is then purified to afford about 50mg of the final compound.

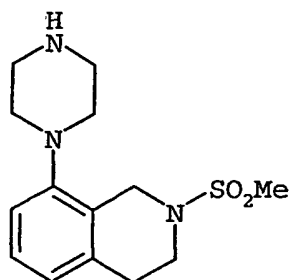
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Preparation 55A

5-(4-t-Butyloxycarbonylpiperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline is treated to the methyl sulfonylation and then deprotection as described in Steps 2 and 3 of
25 Preparation 70A to afford about 45 mg of the final compound (80%).

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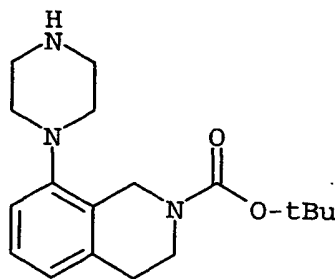
Preparation 56A

Step 1: 8-(Piperazin-1-yl)isoquinoline is prepared from 8-bromoisquinoline and piperazine using the coupling procedure as described in Step 1 of Preparation 54A to afford about 270mg (88%). EIS-MS 214.0 M+1

10 Step 2: 8-(Piperazin-1-yl)isoquinoline (250mg, 1.17mmol) is dissolved in DCM (15mL). Boc₂O (1.02g, 4.69mmol) is added in one portion. The mixture is stirred for about 2 hours at r.t. and then concentrated. The residue is chromatographed on silica gel to give about 327 mg of 8-(4-t-butyloxycarbonyl-piperazin-1-yl)isoquinoline (89%).

15 Step 3: The compound of Step 2 (2.8g, 8.93mmol) is dissolved in isopropanol (120mL), and platinum oxide (1g) is added. The mixture is pressurized to 45psi under hydrogen and allowed to react for about 12 hours. The mixture is filtered, concentrated and chromatographed to give about 1.98g of 8-(4-t-butyloxycarbonyl-piperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline (70%).

20 Step 4: The compound of Step 3 is treated to the methyl sulfonylation and deprotection as described in Preparation 70A to afford about 100mg of the final compound (77%). EIS-MS 296.1 M+1

Preparation 57A

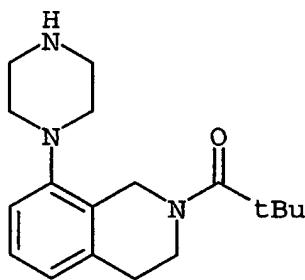
-115-

5 Step 1: 8-(Piperazin-1-yl)isoquinoline (250mg, 1.17mmol) is placed in DCM (5mL) with pyridine (283 microliters, 3.51mmol), and trifluoroacetic anhydride (182 microliters, 1.29mmol) is added dropwise for about 5 minutes. The mixture is stirred at r.t. for about 4 hours and then diluted with DCM and washed with water. The water is extracted with diethyl ether, and the organic extracts are combined, dried, filtered
10 and concentrated to afford a crude mixture containing 8-(4-trifluoroacetyl-piperazin-1-yl)isoquinoline, which is carried forward without further purification. EIS-MS 310.3 M+1

Step 2: The mixture from Step 1 is dissolved in isopropanol (75mL), and Pt/C 10% (200mg) is added. The mixture is pressurized to 45psi for about 12 hours and
15 then filtered and concentrated. The residue is purified via chromatography on silica gel to give about 35mg of 8-(4-trifluoroacetyl-piperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline.

Step 3: The compound of Step 2 (35mg, 0.14mmol) is dissolved in DCM (5mL), and Boc₂O (46mg, 0.14mmol) is added. The mixture is stirred under nitrogen at r.t. for about 2 hours. The mixture is then concentrated in vacuo and plug
20 filtered on silica gel. The filtrate is dissolved in a 2M methanolic solution of ammonia (5mL) and stirred at r.t. under nitrogen overnight. The mixture is evaporated to dryness to afford about 45 mg of the final product.

Preparation 58A

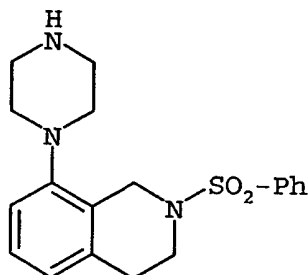


25 8-(4-t-butyloxycarbonyl-piperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline (150mg, 0.47mmol) is dissolved in DCM (5mL), and Et₃N (786 microliters, 3.76mmol) is added. To the stirred system is added pivaloyl chloride (232 microliters, 1.88mmol) by syringe under nitrogen. The mixture is stirred for about 3 hours at r.t. and then
30 concentrated under vacuum. Chromatography on silica gel afforded the Boc-protected

-116-

- 5 compound, which is dissolved in DCM (2.5mL) and TFA (2.5mL) and stirred at r.t. for about 3 hours. The solution is concentrated and the residue is purified by SCX chromatography to afford about 44 mg of the final compound (31%).

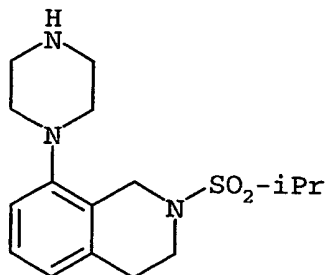
Preparation 59A



- 10 8-(4-t-butylloxycarbonyl-piperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline (127mg, 0.40mmol) is dissolved in DCM (5mL), and Et₃N (446 microliters, 3.2mmol) is added. To the stirred mixture is added phenyl sulfonyl chloride (204 microliters, 1.6mmol) by syringe under nitrogen. The mixture is stirred for about 4 hours and then
- 15 concentrated under vacuum. Chromatography on silica gel affords the Boc-protected product. The protected product is taken up in DCM (2.5mL) and TFA (2.5mL) and the resulting mixture is stirred for about 2 hours. The mixture is concentrated and the residue is purified by SCX chromatography to afford about 119mg of the final compound (83%).
- EIS-MS 358.1 [M+1]

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Preparation 60A



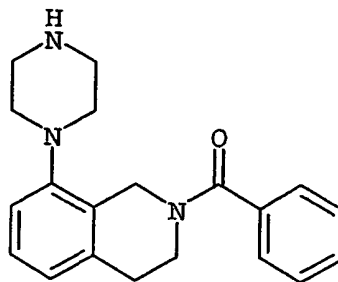
- 25 8-(4-t-butylloxycarbonyl-piperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline (127mg, 0.40mmol) is dissolved in DCM (5mL), and Et₃N (446 microliters, 3.2mmol) added. To the stirred solution is added isopropyl sulfonyl chloride (180 microliters,

-117-

- 5 0.6mmol) by syringe under nitrogen. The mixture is then stirred for about 4 hours and concentrated under vacuum. Chromatography on silica gel afforded the Boc-protected product, which is taken up in DCM (2.5mL) and TFA (2.5mL) and stirred for about 3 hours. The solution is concentrated and the residue is purified by SCX chromatography to afford about 57mg of the final compound (44%). EIS-MS 322.1 [M+1]

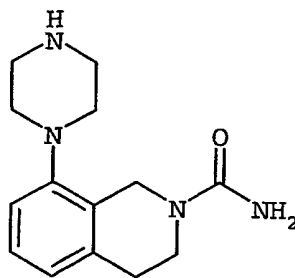
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Preparation 61A

8-(4-t-butyloxycarbonyl-piperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline (127mg, 0.40mmol) is dissolved in DCM (5mL), and Et₃N (446 microliters, 3.2mmol) added. To the stirred solution is added benzoyl chloride (186 microliters, 1.6mmol) by syringe under nitrogen. The mixture is then stirred for about 4 hours and concentrated under vacuum. Chromatography on silica gel afforded the Boc-protected product, which is taken up in DCM (2.5mL) and TFA (2.5mL) and stirred for about 3 hours. The mixture is concentrated and the residue is purified by SCX chromatography to afford about 104mg of the final compound (82%). EIS-MS 322.1 (M+1).

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Preparation 62A

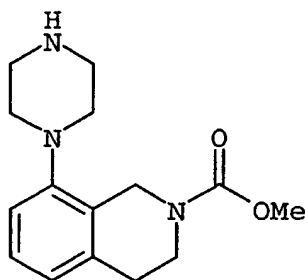
8-(4-t-butyloxycarbonyl-piperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline (250mg, 0.79mmol) is dissolved in acetonitrile (5mL) and DMF (2mL). To the stirred solution is added sodium cyanate (103mg, 1.58mmol). The mixture is heated to 75°C for about 2 hours and allowed to cool to r.t., which is then stirred for an additional 48 hours. After adding TFA (122 microliters, 1.58mmol), the mixture is stirred for another hour, which is then concentrated and chromatographed. The resulting product is dissolved in DCM (2mL) and TFA (2mL) and stirred for about an hour. The product is concentrated

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-119-

- 5 and purified by SCX chromatography to afford about 113mg of the final compound (55%). EIS-MS 261.1 (M+1)

Preparation 63A

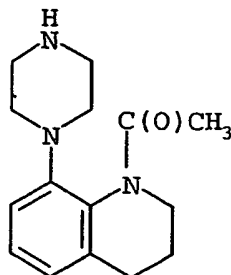


- 10 8-(4-t-butyloxycarbonyl-piperazin-1-yl)-1,2,3,4-tetrahydroisoquinoline (250mg, 0.79mmol) is dissolved in DCM (5mL), and Et₃N (661 microliters, 4.74mmol) is added. To the stirred system is added methyl chloroformate (183 microliters, 2.37mmol) by syringe under nitrogen. The mixture is stirred for about 2 hours and then concentrated under vacuum. The residue is partitioned between DCM and aqueous
- 15 sodium bicarbonate solution. Chromatography on silica gel of the concentrated organic layer afforded the Boc-protected product, which is taken up in DCM (2.5mL) and TFA (2.5mL) and stirred for about 3 hours. The product is concentrated and purified by SCX chromatography to afford about 113mg of the final compound (52%). EIS-MS 276.1 [M+1]

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Preparation 64A

Step 1: 8-(Piperazin-1-yl)quinoline is prepared from 8-bromoquinoline and piperazine as described in Step 1 of Preparation 54A to afford about 490mg (95%). EIS-MS 207.8 (M+1).

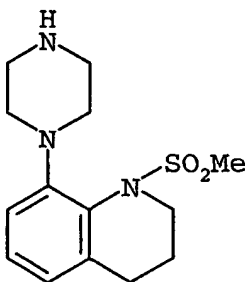
10 Step 2: 8-(Piperazin-1-yl)quinoline (490mg, 2.3mmol) is dissolved in DCM (30mL), and Boc₂O (2g, 9.2 mmol) is added in one portion. The mixture is stirred for about 2 hours at r.t. and then concentrated. The residue is purified to give about 644mg of 8-(4-t-butyloxycarbonylpiperazin-1-yl)quinoline (89%).

15 Step 3: The compound of Step 2 (280mg, 0.89mmol) is dissolved in EtOH (70mL), and Pt/C 10% (300 mg) is added. The mixture is pressurized to 45psi under hydrogen and reacted for about 4 hours. The mixture is filtered over celite and concentrated. The residue is purified to afford about 266 mg of 8-(4-t-butyloxycarbonylpiperazin-1-yl)-1,2,3,4-tetrahydroquinoline (94%). EIS-MS 318.1 [M+1]

20 Step 4: The compound of Step 3 (300mg, 0.95mmol) is dissolved in acetic anhydride (5mL) and stirred for about 3 hours at r.t. The mixture is diluted with water and DCM (50mL/50mL) and then sodium hydroxide (5M) is added. The organic fraction is concentrated, and the residue is taken up in DCM (2.5mL) and TFA (2.5mL), which is then stirred for about an hour. The mixture is concentrated and purified by SCX
25 chromatography to afford about 33mg of the final compound (13%). EIS-MS 260.2 [M+1]

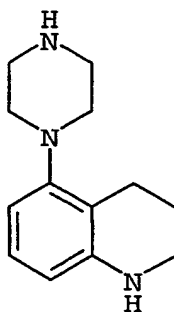
-121-

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Preparation 65A

8-(4-t-butyloxycarbonylpiperazin-1-yl)-1,2,3,4-tetrahydroquinoline (200mg, 0.63mmol) is dissolved in pyridine (5mL), and methane sulfonyl chloride (1.86mL, 12.61mmol) is added. The mixture is stirred for about 4 hours at r.t., which is then concentrated under vacuum and purified via silica gel chromatography. The Boc-protected product is dissolved in DCM (2.5mL) and TFA (2.5mL) and then stirred for about 1.5 hours. The product is concentrated and purified by SCX chromatography to afford about 58mg of the final compound (31%). EIS-MS: Found 296.1 [M+1]

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Preparation 66A

5-Aminoquinoline (5g, 34.7mmol) is dissolved in HBr 48% (15mL) and water (15mL) and then cooled to 0°C. A solution of NaNO₂ (2.5g, 36.2mmol) in water (15mL) is added over about 10 minutes. The mixture is stirred for about 30 minutes at 0°C and then at r.t. for about 1.5 hours. The mixture is transferred to an addition funnel and 75°C solution of CuBr is added over 30 minutes. The mixture is stirred for about 30 minutes and then allowed to cool to r.t., which is then stirred for another 2 hours. The mixture is cooled to 0°C and adjusted to pH of about 14 with 5N NaOH. The solution is filtered, and the solids are washed with DCM. The aqueous layer is extracted with DCM

-122-

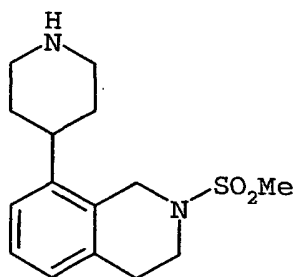
5 (2X 200mL), and the organic fractions are combined and concentrated. The residue is chromatographed on silica gel to give about 1.6g of 5-bromoquinoline (22%).

5-Bromoquinoline is converted to 5-(piperazin-1-yl)quinoline by coupling procedure as described in Step 1 of Preparation 54A to afford about 430mg of the product (84%).

10 5-(piperazin-1-yl)quinoline (430mg, 1.98mmol) is placed in DCM (25mL), and Boc₂O (1.73g, 7.92mmol) is added in one portion. The mixture is stirred for about 2 hours, and then concentrated and purified by SCX chromatography. The Boc-protected product is dissolved in isopropanol (75mL), and Pt/C 5% (400mg) is added. The mixture is pressurized to 45psi under hydrogen and shaken for about 4 hours. The mixture is
15 filtered through celite and the product is isolated via chromatography. The material is placed in DCM (10mL) and TFA (1mL), and then stirred for about 2 hours. The product is concentrated and purified by SCX chromatography to afford about 45mg of the final compound (11%).

20

Preparation 67A



Step 1: 8-Bromoisoquinoline (300mg, 1.44mmol) is dissolved in DMF (8.7mL). To the solution is added Boc-piperidine boronic ester (405mg, 1.31mmol), dichloro(1,1'-bis(diphenylphosphino)ferrocene)palladium (II) dichloromethane adduct (58mg, 0.079mmol) and potassium carbonate (544mg, 3.93mmol). The mixture is heated
25 to about 90°C overnight and cooled, which is then diluted with DCM and filtered through celite. The filtrate is concentrated under vacuum to remove DCM and partitioned between EtOAc (100mL), hexanes (100mL) and water (100mL). The organic fraction is concentrated and chromatographed on silica gel to give about 298mg of 8-(1-t-

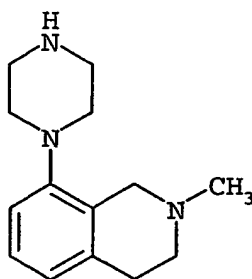
-123-

- 5 butyloxycarbonyl-1,2,5,6-tetrahydropyridin-4-yl)isoquinoline (73%). EIS-MS 311.2 M+1

Step 2: The compound of Step 1 (298mg, 0.96mmol) is dissolved in isopropanol (100mL), and platinum oxide is added (500mg). The mixture is pressurized to 45psi under hydrogen and shaken overnight. The mixture is filtered, concentrated and
10 chromatographed on silica gel to give about 232mg of 8-(1-t-butyloxypiperidin-4-yl)-1,2,3,4-tetrahydroisoquinoline (76%). EIS-MS 317.2 M+1

Step 3: The compound of Step 2 (232mg, 0.73mmol) is treated to the methyl sulfonylation and then deprotected as described in Preparation 70A. The crude product is concentrated and purified by SCX chromatography to afford about 125mg of
15 the final product (58%). EIS-MS 295.1 (M+1)

Preparation 68A



- 8-Bromoisoquinoline and 1-Boc-piperazine are coupled as described in
20 Step 1 of Preparation 54A. Formaldehyde (0.095ml, 1.2mmol, 38% aqueous solution), sodium triacetoxyborohydride (374mg, 1.8mmol) in glacial acetic acid (0.23ml, 3.8mmol) and 1,2-dichloroethane (3ml) are added to a slurry of the coupled compound (200mg, 0.63mmol). The mixture is stirred for about 16 hours at r.t. under nitrogen, and then 1N NaOH is added. The mixture is diluted with ether (25ml) and then washed with brine.
25 The ether extract is concentrated under vacuum to give oil. The oil is dissolved in 3ml of DCM and 1ml of TFA. The mixture is stirred at r.t. overnight, and then concentrated under vacuum and loaded onto a 10 gram SCX (cation exchange resin). The product is eluted with 2M NH₃ in methanol and concentrated under vacuum to afford about 138mg of the final compound (89%).

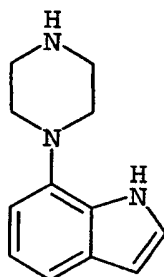
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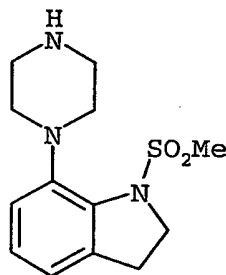
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Preparation 69A

7-(Piperazin-1-yl)indole



The Boc protected indole (200mg, 1.5mmol) prepared from 7-bromoindole and N-Boc piperazine as described in Step 1 of Preparation 54A is dissolved in DCM
10 (1mL). TFA (1mL) is added and the mixture is stirred at r.t. for about an hour. The mixture is then concentrated to oil, which is taken up in methanol and freebased via SCX purification to afford about 164 mg (54%) of the final compound.

Preparation 70A

15

Step 1: 7-(4-t-Butyloxycarbonylpiperazin-1-yl)indole (200mg, 0.67mmol) is dissolved in acetic acid (4mL). To the solution, sodium cyanoborohydride (84 mg, 1.33mmol) is added. The reaction mixture is stirred for about 2 hours at r.t. The mixture is then diluted with DCM and NaOH (1M) is added. The organic layer is separated, dried
20 and concentrated. The residue is chromatographed on the silica gel and the product containing fractions are combined to give about 93mg of the 7-(4-t-butylloxycarbonylpiperazin-1-yl)-2,3-dihydroindole (46%).

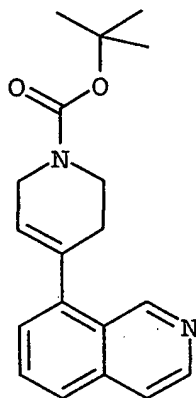
Step 2: To the compound of Step 1 (94mg, 0.31mmol) dissolved in DCM (5mL), TEA (259 microliters, 1.86mmol) is added while stirring. Methane sulfonyl
25 chloride (91 microliters, 0.62mmol) is added by syringe under nitrogen and the mixture is

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5 stirred for about an hour at r.t. The mixture is concentrated under vacuum, and the residue is purified on silica gel to give the Boc protected compound (85mg, 71%).

Step 3: The compound of Step 3 is dissolved in a 1:1 TFA/DCM solution (4mL) for deprotection, and the mixture is stirred for about 1.5 hour at r.t. The product is concentrated to oil, which is then subjected to SCX purification to afford about 53mg of
10 the final compound (85%).

Preparation 71A

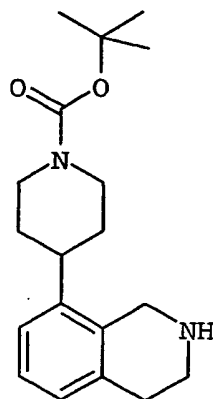


The 1-2(H)-pyridine-carboxylic acid, 3,6-dihydro-4-(4,4,5,5-tetramethyl-
15 1,3,2-dioxaborolan-2-yl)-1,1-dimethyl ethyl ester (1.35 g, 4.37 mmol, *Tet. Lett.*, 41:3705-3708, 2000), 8-bromoisoquinoline (1 g, 4.8 mmol), dichloro(1,1'-bis(diphenylphosphino)ferrocene) palladium (II) DCM adduct (193 mg, 0.263 mmol) and potassium carbonate (1.8 g, 13.1 mmol) are dissolved in DMF (29 mL), degassed and heated at 90°C for about 4 hours.
20 The reaction is cooled to r.t., diluted with DCM (100 mL) and filtered over celite. The filtrate is concentrated to dryness. The resulting residue is partitioned between 600mL of 1:1 EtOAc/hexanes and water (200 mL). The organic phase is concentrated and purified by silica gel chromatography to give about 1.07g of the final product (79%).

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Preparation 72A

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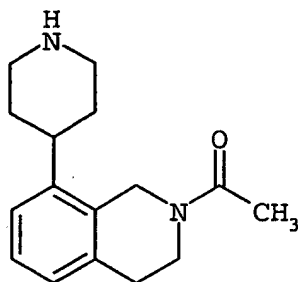
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The compound 71A (1.05 g, 3.38 mmol) is dissolved in a slurry of PtO_2 (1 g) in isopropanol (150 mL), and pressurized to 50 psi with hydrogen overnight. The mixture is filtered, concentrated and purified by silica gel chromatography to give about 1.07g of the final product. EIS-MS 317.2 [M+1]

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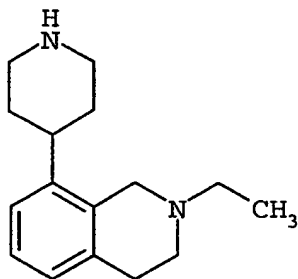
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Preparation 73A

The compound 72A (530 mg, 1.73 mmol) is dissolved in acetic anhydride (10 mL) and TEA (3 mL) is added. The mixture is stirred for about 16 hours, and then concentrated and dissolved in a 1:1 mixture of DCM/TFA (10 mL). The mixture is stirred for about 2 hours and then concentrated. The resulting residue is subjected to SCX ion exchange chromatography followed by silica chromatography to give about 218 mg of the final product (49%). EIS-MS 259.1 [M+1]

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Preparation 74A

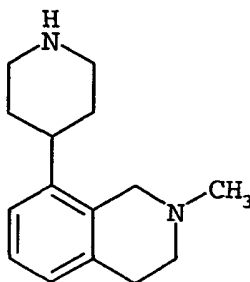
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The compound 73A (287 mg, 1.11 mmol) is dissolved in THF (10 mL) and lithium aluminum hydride (169 mg, 4.44 mmol) is added as a slurry in THF (5 mL). The mixture is stirred for about 4 hours at reflux. The reaction is worked up with water, washed with 1M NaOH and purified by silica gel chromatography to give about 215mg of the final product (79%). EIS-MS 245.2 [M+1]

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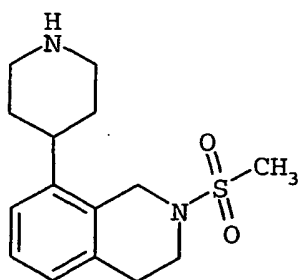
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Preparation 75A

The compound 72A (530 mg, 1.67 mmol) is dissolved in dichloromethane (10 mL) to which is added formaldehyde 38% (8.8 mL, 3.34 mmol), sodium triacetoxyborohydride (1.06 g, 5.02 mmol) and acetic acid (0.6 mL). The mixture is stirred at r.t. for 24 hours. The mixture is diluted with DCM (50 mL) and washed with 1M NaOH. The organic fraction is concentrated and purified by silica gel chromatography. The product is then stirred in 1:1 TFA/DCM (10 mL) for about 2 hours. The reaction is concentrated, and the resulting residue is subjected to SCX ion exchange chromatography followed by silica gel chromatography to give about 171mg of the final product (44%). EIS-MS 231.2 [M+1]

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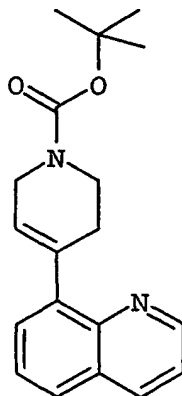
Preparation 76A

To the compound 72A (225 mg, 0.71 mmol) in DCM (10 mL) is added TEA (1.2 mL, 8.6 mmol) and methane sulfonyl chloride (0.329 mL, 4.3 mmol), and the mixture is stirred at r.t. for about 3 hours. The mixture is diluted with DCM (50 mL) and washed with 1M NaOH. The organic fraction is concentrated and the resulting residue is stirred in 1:1 TFA/DCM (10 mL) for about 2 hours. The mixture is concentrated, and the residue is subjected to SCX ion exchange chromatography followed by silica gel chromatography to give about 125 mg of the final product (60%). EIS-MS 295.0 [M+1]

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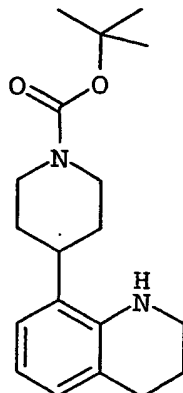
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Preparation 77A

- To 1-2(H)-pyridine-carboxylic acid, 3,6-dihydro-4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1,1-dimethyl ethyl ester (500 mg, 1.62 mmol, *Tet. Lett.*, 41:3705-3708, 2000) in DMF (9 mL) is added 8-bromoquinoline (370 mg, 1.78 mmol), dichloro(1,1'-bis (diphenylphosphino)ferrocene) palladium (II) DCM adduct (71 mg, 0.097 mmol) and potassium carbonate (672 mg, 4.86 mmol), and the mixture is degassed and heated at 90°C for about 4 hours. The mixture is then cooled to r.t., diluted with DCM (100 mL) and filtered over celite. The filtrate is concentrated to dryness. The mixture is partitioned between 1:1 EtOAc/hexanes (600 mL) and water (200 mL). The organic phase is concentrated to dryness, and the resulting residue purified by flash chromatography to give about 230mg of the final product (74%). EIS-MS 311.1 [M+1]

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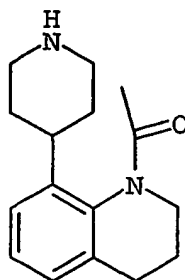
Preparation 78A

To the compound 77A (270 mg, 0.87 mmol) in acetic acid (4 mL) is added sodium cyanoborohydride (110 mg, 1.75 mmol), and the mixture is stirred for about 4 hours. The mixture is diluted with DCM (100 mL) and made basic with 5N NaOH solution. The organic phase is dried and concentrated, and the resulting residue is dissolved in a slurry of 5%Pt/C (500 mg) in isopropanol (150 mL). The mixture is pressurized to 50 psi with hydrogen overnight. The mixture is filtered, concentrated and purified by silica gel chromatography to give about 175mg of the final product (65%).

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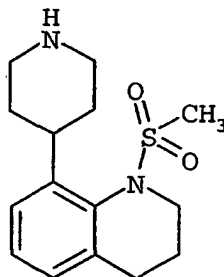
-131-

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Preparation 79A

To compound 78A (175 mg, 0.55 mmol) is added acetic anhydride (10 mL) and pyridine (3 mL), and the mixture is stirred for about 16 hours. After concentrating to dryness, the resulting residue is dissolved in a 1:1 mixture of DCM/TFA (10 mL) and stirred for about 2 hours, which is then concentrated to dryness. The resulting residue is subjected to SCX ion exchange chromatography followed by silica gel chromatography to give about 39mg of the final product (27%). EIS-MS 259.1 [M+1]

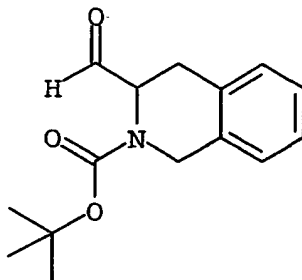
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Preparation 80A

To the compound 78A (555 mg, 1.58 mmol) in pyridine (15 mL) is added methane sulfonyl chloride (2 mL). The mixture is stirred for about 16 hours at r.t. and concentrated, which is then dissolved in a 1:1 mixture of DCM/TFA (10 mL). The mixture is stirred for about 2 hours. After concentrating, the resulting residue is subjected to SCX ion exchange chromatography followed by silica chromatography to give about 218mg of the final product (47%). EIS-MS 295.1 [M+1]

Preparation of "C Domain"Preparation 1C

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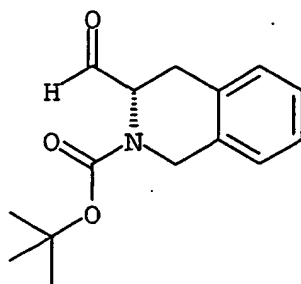
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Boc-D-Tic-OH (4.50g, 16mmol), methoxymethylamine hydrochloride (1.56g, 16mmol), HOBT (2.43g, 18mmol), EDC (3.45g, 18mmol), DMAP (200mg, 1.6mmol), DIPEA (4.2ml) and THF (150ml) are combined, and the resulting mixture is stirred for about 18 hours at r.t. The mixture is stripped to dryness, and the resulting residue is taken up in ethyl acetate and then washed with aqueous HCl (1N), aqueous sodium bicarbonate and brine, which is then dried over sodium sulfate. The solvent is removed and the residue is chromatographed on normal phase (ethylacetate/hexanes 1:1) to give about 3.89g of Boc-D-Tic-NMeOMe (Weinreb amide).

Lithium aluminum hydride (1.0M in THF, 7.2ml, 7.19mmol) is slowly added to the Weinreb amide (1.84g, 5.75mmol) in THF (50ml) at 0°C. The mixture is stirred at 0°C for another 15 minutes. Aqueous KHSO₄ (1.37g in 29ml H₂O) is slowly added followed by diethyl ether. The organic layer is separated, and the aqueous layer is extracted with diethyl ether. The organic phases are combined and washed with aqueous 1M HCl, saturated aqueous NaHCO₃ solution and brine, which is then dried over Na₂SO₄. Removal of solvent affords about 1.44g of the epimerized compound. EIS-MS: MH⁺ 262 [M+1]

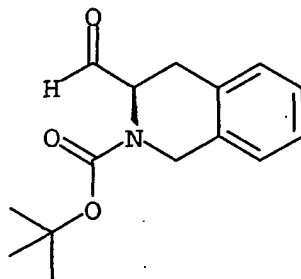
-133-

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Preparation 2C

The above compound was prepared from Boc-L-Tic-OH as described in Preparation 3C below, except that the Weinreb amide was made by a similar procedure to that described in *Synthesis*, 676, 1983.

10

Preparation 3C

Boc-D-Tic-OH (14.9g, 53.7mmol), methoxymethylamine hydrochloride (5.24g, 53.7mmol), EDC (11.3g, 59.1mmol), HOBT (7.98g, 59.1mmol), DIPEA (9.83ml, 59.1mmol) and THF (500ml) are combined, and the resulting mixture is stirred for about 18 hours at r.t. under nitrogen. The mixture is stripped to dryness, and the residue is taken up in ethyl acetate. The resulting mixture is washed with 1M HCl, saturated NaHCO₃ and brine, and then dried. Solvent is removed and the residue is chromatographed on normal phase using 1:1 ethylacetate/hexane to give about 12.3g of Boc-D-Tic-NMeOMe (Weinreb amide).

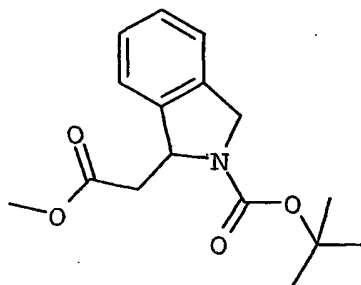
Lithium aluminum hydride (1.0M in THF, 5.1ml, 5.00mmol) is slowly added to the Weinreb amide (1.28g, 4.00mmol) in THF (35ml) at 0°C. The mixture is stirred at 0°C for another 15 minutes. Aqueous KHSO₄ (970mg in 20ml H₂O) is slowly added followed by diethyl ether. The organic layer is separated, and the aqueous layer is extracted with diethylether. The organic phases are combined and washed with aqueous

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- 5 1M HCl, saturated aqueous NaHCO_3 and brine, which is then dried over Na_2SO_4 .
Removal of solvent affords about 780mg of the final compound. EIS-MS: MH^+ 262
[M+1]

Preparation 4C

- 10 1-Methoxycarbonylmethyl-1,3-dihydro-isoindole-2-carboxylic acid tert-butyl ester



Step A: (2-Bromo-benzyl)-carbamic acid tert-butyl ester

- To a mixture of 125.0 g (561.8 mmol) of 2-bromobenzylamine hydrochloride and 170.7 g
(1236.0 mmol) of potassium carbonate in 300 mL of 50% THF/water is added 134.9 g
15 (618.0 mmol) of di-*tert*-butyl dicarbonate in four portions over 20 minutes. The mixture
is stirred at r.t. for about 16 hours and then diluted with 300 mL of ethyl acetate and 300
mL of water. The organic portion is separated and the aqueous portion is extracted three
times with 200 mL each of ethyl acetate. The combined ethyl acetate portions are washed
once with 250 mL of 10% aqueous sodium bisulfate. The organic portion is dried
20 (MgSO_4), filtered and concentrated to dryness to afford about 161g Step A compound.

Step B: 3-[2-(tert-Butoxycarbonylamino-methyl)-phenyl]-acrylic acid
methyl ester

- To compound of Step A (161.0 g, 561.8 mmol) in DMF (800 mL) is added methyl
acrylate (58.0 g, 674.2 mmol), TEA (170.5 g, 1685.4 mmol) and
25 dichlorobis(triphenylphosphine)
palladium(II) (7.9 g, 11.2 mmol). The mixture is heated at 80°C for about 32 hours. The
mixture is cooled, diluted with 1000 mL of EtOAc and washed with 10% aqueous sodium
bisulfate. The aqueous portion is extracted three times with EtOAc and the combined
organics are dried (Na_2SO_4) and concentrated to dryness. The residue is dissolved in a
30 small amount of DCM and filtered through 7 inch of silica gel in a 2 L sintered glass

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- 5 funnel eluting with 25% EtOAc/hexanes. The eluent is concentrated to dryness and recrystallized from EtOAc/hexanes to afford about 116.9 g (71%) of Step B compound.

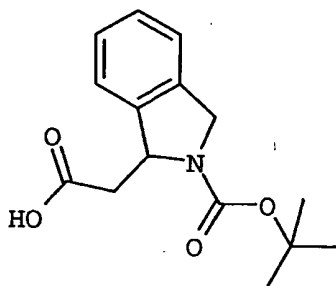
Step C: To a 0°C solution of (116.9 g, 401.2 mmol) material from Step B in DCM (800 mL) is added 200 mL of TFA dropwise over 15 minutes. After removing the cooling bath, the mixture is stirred for about 2.5 hours and then concentrated to dryness. The residue is dissolved in 500 mL of DCM and saturated aqueous sodium bicarbonate is slowly added until the mixture is slightly basic. The organic portion is separated and the aqueous portion is extracted two times with DCM. The combined organic portions are dried (Na₂SO₄) and concentrated to dryness. The residue is dissolved in 800 mL of DCM and DIPEA (57.0g, 441.4 mmol) is added. To the mixture is added di-*tert*-butyl dicarbonate (96.3 g, 441.4 mmol) in five portions over 45 minutes and then stirred at room temperature for 16 h. The mixture is washed with 10% aqueous sodium bisulfate, and the organic portion is separated and the aqueous portion is extracted two times with DCM. The combined organic extracts are dried ((Na₂SO₄) and concentrated to dryness. The resulting residue is dissolved in a small amount of DCM and filtered through 7 inch silica gel in a 2L sintered glass funnel eluting with 25% EtOAc/hexanes. The eluent is concentrated to dryness and the enantiomers are separated by chiral chromatography. The first eluting isomer is labeled as isomer #1 and the second eluting is labeled as isomer #2, which affords about 52.6g (45%) of the final compound (isomer 2). EIS-MS 292 [M+1]

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Preparation 5C

1-Carboxymethyl-1,3-dihydro-isoindole-2-carboxylic acid tert-butyl ester

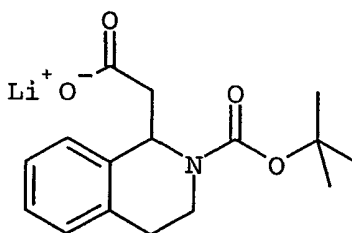


To the compound 4C (52.6 g, 180.5 mmol) in MeOH (500 mL) is added 1 N NaOH (199 mL, 199.0 mmol). The mixture is stirred at r.t. for about 48 hours and then concentrated to dryness. The resulting residue is dissolved in water (300 mL) and extracted with diethyl ether (2x). The aqueous portion is acidified to pH 2 with 10% aqueous sodium bisulfate and extracted with EtOAc. The combined organic extracts are dried (MgSO₄) and concentrated to dryness to afford about 49.8g of the final compound (99%). EIS-MS 276 [M-1].

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Preparation 6C

1-Carboxymethyl-3,4-dihydro-1H-isoquinoline-2-carboxylic acid tert-butyl ester, lithium salt



20

Step A: (1,2,3,4-Tetrahydro-isoquinolin-1-yl)-acetic acid methyl ester

To Boc-tetrahydroisoquinoline-1-acetic acid (100.4 g, 520.0 mmol) in MeOH (200 mL) is added 400mL of 2.3 M HCl in methanol. The mixture is stirred overnight and then concentrated to dryness. The resulting residue is dissolved in EtOAc and washed with saturated sodium bicarbonate and brine, which is then dried (Na₂SO₄) and concentrated to dryness to afford about 109.5g (100%) of the ester. EIS-MS 206 [M+1]

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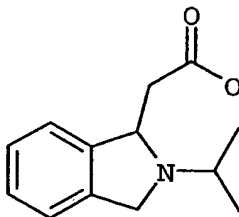
-137-

5 Step B: 1-Methoxycarbonylmethyl-3,4-dihydro-1H-isoquinoline-2-carboxylic acid tert-butyl ester

To a 0°C solution of compound from Step A (50.5g, 240.0 mmol) in THF (250 mL) is added di-*tert*-butyl dicarbonate (59.3 g, 270.0 mmol) in THF (50 mL) dropwise. After stirring for about 45 minutes, the mixture is concentrated to dryness. The resulting
10 residue is dissolved in EtOAc and washed with saturated sodium bicarbonate and brine, which is then dried (Na₂SO₄) and concentrated to dryness. Chiral chromatography of the residue affords both enantiomers, the first eluting isomer being labeled as isomer 1 and the second as isomer 2. EIS-MS 306 [M+1]

Step C: To a solution of compound B from Step B (10.2g, 33.4 mmol) in
15 dioxane (220 mL) is added a solution of lithium hydroxide monohydrate (1.67 g, 39.8 mmol) in water (110 mL) portion wise while maintaining a temperature below 30°C. The mixture is stirred for about 16 hours and then concentrated to dryness, which gives about 11.2g of the final compound. EIS-MS 292 [M+1]

20 Preparation 7C
(2-isopropyl-2,3-dihydro-1H-isoindol-1-yl)-acetic acid



Step A: (2,3-dihydro-1H-isoindol-1-yl)-acetic acid methyl ester:

25 To the compound prepared in Preparation C1 (11.75 g., 40.41 mmol) in DCM (50 mL) was added TFA (50 mL) dropwise. After about 2 hours, the mixture was concentrated to dryness and the resulting residue was partitioned with saturated aqueous sodium bicarbonate (200 mL) and EtOAc (300 mL). The organic portion was separated and the aqueous layer was extracted with DCM (4 x 500 mL). The combined DCM extracts were
30 combined, dried (Na₂SO₄), and concentrated to dryness to afford about 3.97 g (51%).

Step B: (2-isopropyl-2,3-dihydro-1H-isoindol-1-yl)-acetic acid methyl ester:

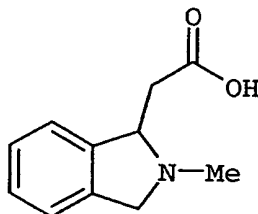
-138-

5 To the compound obtained from Step A (0.50 g, 2.61 mmol) in dichloroethane (46 mL) was added acetone (1.76 mL, 24.01 mmol) and sodium triacetoxyborohydride (2.48 g., 11.74 mmol). After 6 hours, the mixture was diluted with 1.0N NaOH (100 mL), and the organic portion was separated. The aqueous layer was extracted with DCM (3 x 100 mL). The combined DCM extracts were dried (MgSO₄) and concentrated to dryness to afford about 0.60 g (99%). EIS-MS 235 [M+1].

10 Step C: To the compound of Step B (0.53 g., 2.30 mmol) in MeOH (5.1 mL) was added 1.0N NaOH (2.53 mL, 2.53 mmol). After two days, the solution was concentrated to dryness. The resulting residue was diluted with 1.0N HCl and water was loaded onto a strong cation exchange resin. The resin was washed with water, THF/water (1:1) and then water. The product was then eluted from the resin with pyridine/water (1:9). The eluent was concentrated to dryness to afford about 0.43 g (85%) of the final compound. EIS-MS 220 [M+1].

Preparation 8C

20 (2-Methyl-2,3-dihydro-1H-isoindol-1-yl)-acetic acid



Step A: (2-Methyl-2,3-dihydro-1H-isoindol-1-yl)-acetic acid methyl ester:

The compound from preparation C1 was deprotected with TFA in a manner similar to preparation 3C of Step A. To the deprotected compound (0.50 g, 2.61 mmol), in dichloroethane (46 mL), was added 37% aqueous formaldehyde solution (1.80 mL, 24.01 mmol) and sodium triacetoxyborohydride (2.48 g., 11.74 mmol). After 3 days, the mixture was diluted with 1.0N NaOH (100 mL). The organic portion was separated and the aqueous layer was extracted with DCM (3 x 100 mL). The combined DCM extracts were dried (Na₂SO₄) and concentrated to dryness. The resulting residue was purified by

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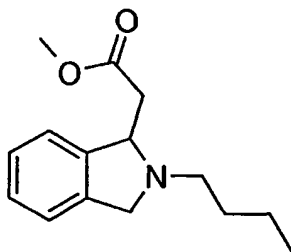
5 flash chromatography (SiO₂, eluting with 100% EtOAc) affording about 0.43 g (79%) of the alkylated isoindole. EIS-MS 206 [M+1].

Step B: To the compound of Step A (0.34 g., 1.66 mmol) in MeOH (3.7 mL) was added 1.0N NaOH (1.82 mL, 1.82 mmol). After 2 days, the solution was concentrated to dryness. The resulting residue was diluted with 1.0N HCl and water was
10 then loaded onto a strong cation exchange resin. The resin was washed with water, THF/water(1:1) and water, and the product was eluted from the resin with pyridine/water(1:9). The eluent was concentrated to dryness to afford about 0.31 g (98%) of the final compound. EIS-MS 192 [M+1].

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Preparation 9C

(2-Butyl-2,3-dihydro-1H-isoindol-1-yl)-acetic acid methyl ester



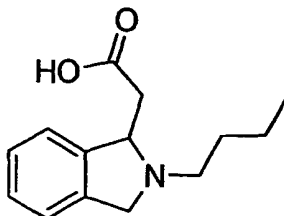
The compound from preparation C1 was deprotected with TFA in a
20 manner similar to preparation 3C of Step A. To the deprotected compound (0.50 g, 2.61 mmol) and butyraldehyde (2.16 mL, 24.01 mmol) in dichloroethane (46 mL) was added sodium triacetoxyborohydride (2.48 g., 11.74 mmol). After reacting about 3 hours, the mixture was diluted with 1.0 N NaOH (100 mL) and partitioned. The aqueous layer was extracted with DCM (3x75 mL). The DCM layers were combined, dried over sodium
25 sulfate, filtered and concentrated under reduced pressure to give a brown residue. The residue was purified via silica gel chromatography (eluent: ethyl acetate/hexanes (1:3). The purified fractions were combined and concentrated to give the title compound as a brown oil (0.51g, 77%). MS ES 249.2 (M+H)

Preparation 10C

30

(2-Butyl-2,3-dihydro-1H-isoindol-1-yl)-acetic acid

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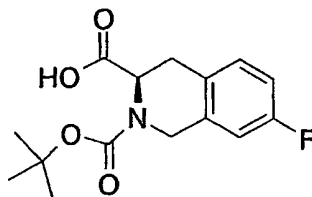


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To a solution containing the compound 7C (0.47 g, 1.89 mmol) in methanol (4.2 mL) was added 1.0 N NaOH (2.08 mL, 2.08 mmol). After reacting about 2 hours, the solution was concentrated under reduced pressure. The residue was diluted with 1.0 N HCl, and water was loaded onto a strong cation exchange resin. The resin was washed with water and THF/water (1:1), and the product was eluted from the resin with pyridine/water (1:9). The pyridine washes were concentrated under reduced pressure, and azeotroped with acetone to give the title compound as brown solids (0.28 g, (64%)) MS ES 234.19 (M+H)

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Preparation 11C



Step A: To a solution of *N*-Boc-4-Fluoro-D-Phe (2.37g, 8.366 mmol) in methanol, 3 mL of concentrated sulfuric acid was added. The mixture was heated to reflux overnight and then concentrated *in vacuo*. MS M+1 198.1

Step B: To an ice cold mixture of 1.65 g (8.367 mmol) of compound from Step A, 1.353 mL of pyridine and ethyl chloroformate (0.848 mL, 8.869 mmol) is added slowly with stirring for about 30 minutes giving white solid. The mixture was partitioned between water and ethyl acetate. The aqueous layer was extracted with EtOAc (2x). The combined organic solution was dried over MgSO₄, filtered, and concentrated *in vacuo* to give about 2.17g of yellow oil (96%). MS M+1 270.1.

Step C: A mixture containing 2.17g (8.06 mmol) of the compound from Step B, paraformaldehyde (0.254g, 8.46 mmol), and 10 mL of 3:1 glacial acetic acid/conc. sulfuric acid was stirred at r.t. for about 48 hours. The mixture was partitioned

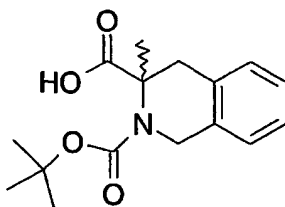
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5 between water and ethyl acetate. The aqueous layer was extracted with EtOAc (3x). The combined EtOAc solution was dried over magnesium sulfate, filtered, and concentrated *in vacuo*. The desired product was purified by column chromatography eluting with 25% EtOAc in Hexane to give about 1.31g (58%) of colorless oil. MS: M+1 282.1

Step D: A solution of 1.31g (4.656 mmol) of material from Step C in 20
10 mL of 5N HCl was heated to reflux for about 24 hours. The solution was concentrated *in vacuo*. The resulting white solid was washed with ether to afford about 0.87g (81%). MS M+1 196.1.

Step E: To a solution of 0.87g (3.755 mmol) of material from Step D in
20 ml of 1:1 dioxane/water, di-*t*-butyl-dicarbonate (0.901g, 4.131 mmol) and 2.355 mL
15 (16.90 mmol) of TEA were added. The mixture was allowed to stir at r.t. overnight. The mixture was diluted with EtOAc, and the separated aqueous layer was extracted with EtOAc (3x). The combined organic solution was dried over magnesium sulfate, filtered and concentrated *in vacuo* to give about 0.64g (58%) of the final compound. MS M-1 294.1.

20

Preparation 12C

Step A: By following a procedure of Preparation 28C, Step A and
25 1.0g(5.58 mmol) of \square -methyl-DL-phenylalanine, about 1.4g of ester was prepared. MS M+1 194.1

Step B: By following a procedure of Preparation 28C, Step B and 1.08g
(5.59 mmol) of material from Step A, about 1.48g (100%) of product was prepared. MS M+1 266.1

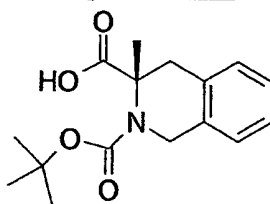
30 Step C: By following a procedure of Preparation 28C, Step C and 1.48g
(5.59 mmol) of material from Step B, about 1.55g (100%) of product was prepared. MS M+1 278.1

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5 Step D: By following a procedure of Preparation 28C, Step D and 1.55g (5.59 mmol) of material from Step C, about 1.33g of product was prepared. MS M+1 192.1

Step E: By following a procedure of Preparation 28C, Step E and 1.33g (5.84 mmol) of material from Step D, about 1.70g (100%) of the final compound was
10 prepared. MS M+1 292.2

Preparation 13C



15 Step A: By following a procedure of Preparation 28C, Step A and 2.0g (11.16 mmol) of α -methyl-D-phenylalanine, about 2.15g of ester was prepared. MS M+1 194.1

Step B: By following a procedure of Preparation 28C, Step B and 2.15g (11.16 mmol) of material from Step A, about 1.46g (49%) of product was prepared. MS
20 M+1 266.1

Step C: By following a procedure of Preparation 28C, Step C and 1.46g (5.503 mmol) of material from Step B, about 0.74g (48%) of product was prepared. MS M+1 278.1

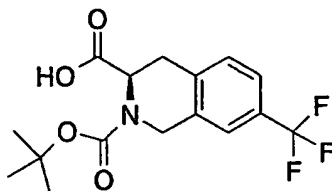
Step D: By following a procedure of Preparation 28C, Step D and 0.74g (2.67 mmol) of material from Step C, about 0.54g (89%) of product was prepared. MS
25 M+1 192.1

Step E: By following a procedure of Preparation 28C, Step E and 0.54g (2.37 mmol) of material from Step D, about 0.54g (78%) of the final compound was prepared. MS M+1 292.2

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Preparation 14C

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Step A: By following a procedure of Preparation 28C, Step A and 0.65g (1.95 mmol) of *N*-Boc-4-trifluoromethyl-D-phenylalanine, about 0.48g of ester was prepared. MS M+1 248.0

10 Step B: By following a procedure of Preparation 28C, Step B and 0.48g (1.95 mmol) of material from Step A, about 0.60g (96%) of product was prepared. MS M+1 320.1

Step C: By following a procedure of Preparation 28C, Step C and 0.6g (1.879 mmol) of material from Step B, about 0.37g (59%) of product was prepared. MS
15 M+1 332.1

Step D: By following a procedure of Preparation 28C, Step D and 0.37g (1.117 mmol) of material from Step C, about 0.11g (35%) of product was prepared. MS M+1 246.1

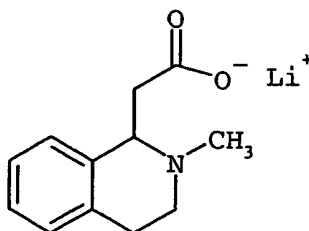
Step E: By following a procedure of Preparation 28C, Step E and 1.11g (0.391 mmol) of material from Step D, about 0.234 g (>100%) of the final compound is
20 prepared. MS M-1 344.1

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Preparation 15C

lithium; (2-methyl-1,2,3,4-tetrahydro-isoquinolin-1-yl)-acetate

Step 1: (1,2,3,4-Tetrahydro-isoquinolin-1-yl)-acetic acid methyl10 ester

The material from Preparation of 13C Step 2 (9.98g, 32.7 mmol) was mixed with 500 mL cold 4M HCl/dioxane and stirred at r.t. for about an hour. The mixture was concentrated *in vacuo*. The residue was dissolved in ethyl acetate and then washed with saturated sodium bicarbonate and brine. The organic portion was dried (Na₂SO₄), filtered and concentrated *in vacuo* to afford about 6.9g (100%) of the title compound. EIS-MS: 206 (M+1).

Step 2: (2-methyl-1,2,3,4-tetrahydro-isoquinolin-1-yl)-acetic acid methyl ester

To a solution of material from Step 1 (6.71g, 32.0mmol) in 175 mL of dichloroethane was added 37% aqueous formaldehyde (22.6mL, 300mmol). After about 10 minute, sodium triacetoxyborohydride (31.2g, 147.0mmol) was added in 2 to 3 g portions with some cooling to maintain ambient temperature. The mixture was stirred for about 16 hours and DCM and water was added. The mixture was adjusted to pH 9-10 with 5N sodium hydroxide. The organic layer was separated, washed with brine, and then dried (Na₂SO₄) and concentrated *in vacuo*. Chromatography (silica gel, 5%(2N ammonia in methanol)/DCM) of the residue afforded about 6.9g (96%) of the title compound. EIS-MS: 220 (M+1).

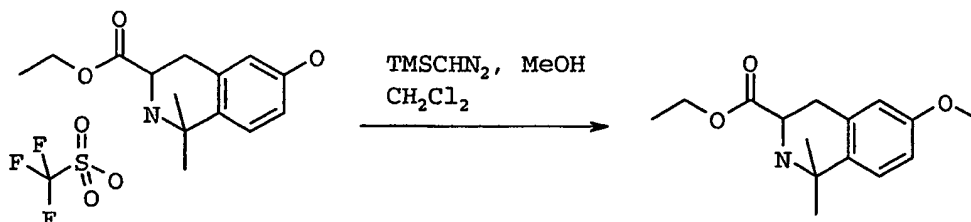
Step 3: To a solution of material from Step 2 (4.45g, 18.9 mmol) in 120 mL dioxane was added lithium hydroxide monohydrate (1.02g, 22.7 mmol) in 65 mL water in portions keeping the temperature below 30°C. After about 16 hours, the mixture was concentrated *in vacuo* to afford about 8.12g of the final compound. EIS-MS: 206 (M+1).

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Preparation 16C

1,1-Dimethyl-6-methoxy-1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid ethyl ester



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To a solution of the triflate salt of 1,1-dimethyl-1,2,3,4-tetrahydroisoquinoline-3-carboxylic acid ethyl ester (1.5 g, 3.76 mmol, 1.0 eq.) in MeOH (20 mL) and CH₂Cl₂ (2 mL) at 0°C was added a solution of (trimethylsilyl)diazomethane (2.0 M in hexane, 3.7 mL, 2.0 eq.). The resulting mixture was warmed to r.t. and stirred overnight, and then the solution was concentrated. Purification by flash chromatography (125 g SiO₂ linear gradient, 40 mL/min, 1:1 EtOAc/hexane for 33 minutes) afforded about 900 mg of the final compound (96%). LRMS (electrospray): 250.2 (M+1).

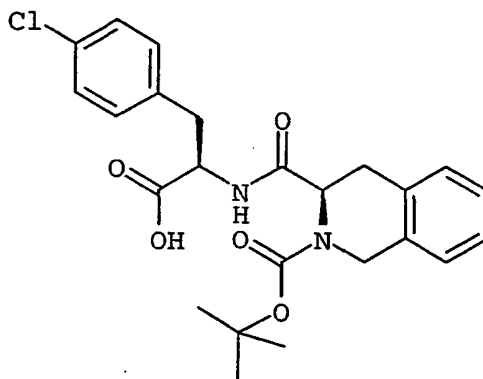
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5 "B Domain" and "C Domain" CombinationPreparation 1BC

3-[2-(4-Chloro-phenyl)-1-methoxycarbonyl-ethyl]carbamoyl]-3,4-dihydro-1H-
isoquinoline-2-carboxylic acid tert-butyl ester
(N-Boc-D-Tic-4-Cl-D-phe-OH)

10



Step A: To a 0°C solution of 4-Cl-D-Phe methyl ester (23.8 g, 111.0 mmol), Boc-D-Tic (30.8 g, 111.0 mmol) and 4-DMAP (75 mg, 0.61 mmol) in 200 mL of DCM is added EDC (30.8 g, 111.0 mmol) and the mixture is stirred for about 20 minutes. The ice bath is removed and the mixture is stirred at r.t. for about 4 hours. After washing with water (4 x 200 mL), the combined aqueous portions are back extracted with DCM (2 x 200 mL). The combined organic portions are washed with brine, dried (MgSO₄) and concentrated to dryness. The desired product is purified by flash chromatography (SiO₂, eluting with 35% EtOAc in Hexanes) to afford about 43.0g (83%) of the ester. EIS MS 473 [M+1]

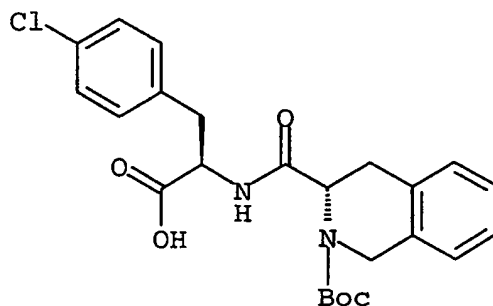
Step B: To the compound of Step A (43.0 g, 91.0 mmol) in MeOH (170 mL) at 0°C is added 1N NaOH (227.0 mL, 227.0 mmol) dropwise. After about 20 minutes, the ice bath was removed and the mixture is stirred at r.t. for about 3 hours. The mixture is concentrated to dryness, and the resulting residue is suspended in 200 mL of water. The aqueous layer is adjusted to about pH 1 with 5 N HCl and then extracted with EtOAc (4 x 200 mL). The combined organics are dried (MgSO₄) and concentrated to dryness yielding about 39.0g (93%) of the final compound. EIS-MS 459 [M+1]

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Preparation 2BC

Boc-L-Tic-4-Cl-D-phe-OH



10 Boc-L-Tic-4-Cl-D-phe-OMe is prepared as described in Preparation 1BC of Step A. $^1\text{H NMR}$ (DMSO) (two rotomers observed) δ 8.40(d, 1H), 8.32(d, 0.5 H), 7.32(d, 2H), 7.25(d, 0.5H), 7.18(d, 2H), 7.00-7.16(m, 4.5H), 6.88(m, 2H), 4.74(m, 0.5H), 4.27-4.60(m, 4.5H), 3.58(s, 3H), 3.56(s, 1.5H), 2.60-3.07(m, 6H), 1.42(s, H), 1.21(s, 9H). MS(ES) 473.0(M^+), 471.1(M).

15 Boc-L-Tic-4-Cl-D-phe-OH is converted to Boc-L-Tic-4-Cl-D-phe-OH-OH according to the procedure described in Preparation 1BC of Step B. $^1\text{H NMR}$ (DMSO) (Two rotomers observed) δ 7.98 (d, 1H), 7.72 (d, 0.5 H), 6.90-7.41 (m, 16H), 4.0-4.70 (m, 8.5H), 2.60- 3.20 (m, 8.5H), 1.32-1.41 (m, 19H). MS(ES) 459.1 m/z(M^+), 457.1(M).

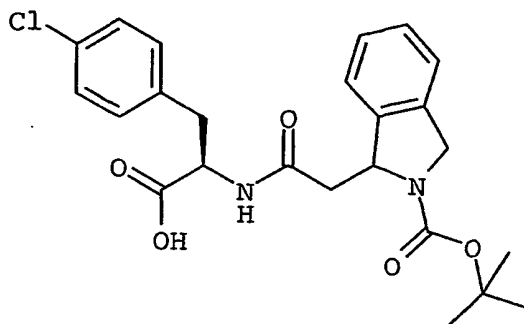
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Preparation 3BC

1-[[1-Carboxy-2-(4-chloro-phenyl)-ethylcarbamoyl]-methyl]-1,3-dihydro-isoindole-2-carboxylic acid tert-butyl ester



Step A: To a suspension of 4-Cl-D-Phe methyl ester hydrochloride (40.4 g, 161.5 mmol) in DCM (250 mL), is added saturated aqueous sodium bicarbonate (250 mL) and the mixture is stirred at r.t. for about one hour. The organic portion is separated and the aqueous portion is extracted with DCM (2x). The combined organic portions are dried (Na_2SO_4) and concentrated to dryness. To the free amine in DCM (400 mL) at 0°C is added the compound 5C (isomer 2, 44.8 g, 161.5 mmol), EDC (31.0 g, 161.5 mmol) and 4-DMAP (2.0 g, 16.1 mmol). The reaction mixture is stirred at 0°C for about 30 minutes and then stirred for another 5 hours at r.t. The mixture is then washed with saturated aqueous sodium bicarbonate (200 mL) and 10% aqueous sodium bisulfate (200 mL). The mixture is dried (Na_2SO_4) and concentrated to afford about 76.4g (100%) of the ester. EIS-MS 471 [M-1].

Step B: To the ester from Step A (76.4 g, 161.5 mmol) in MeOH (760 mL) is added 1 N NaOH (242.0 mL, 242.0 mmol), and the mixture is heated at 50°C for about 4 hours. The mixture is stirred for another 16 hours at r.t. After concentrating to dryness, the resulting residue is taken up in 500 mL of water and washed with diethyl ether (2x). The aqueous portion is acidified to pH 2 with 10% aqueous sodium bisulfate and extracted with EtOAc (4 x 200 mL). The combined organic extracts are dried (MgSO_4) and concentrated to dryness. The resulting solid is suspended in hexane and then filtered and dried to afford about 67.7g (91%) of the final compound. EIS-MS 457 [M-1].

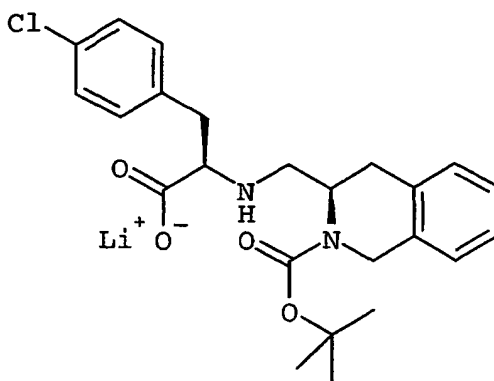
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Preparation 4BC

Lithium; 2-[(2-tert-butoxycarbonyl-1,2,3,4-tetrahydro-isoquinolin-3-ylmethyl)-amino]-3-(4-chloro-phenyl)-propionate



- Step A: 3-(Methoxy-methyl-carbamoyl)-3,4-dihydro-1H isoquinoline-2-
 10 carboxylic acid tert-butyl ester
 To Boc-D-1,2,3,4-tetrahydroisoquinoline carboxylic acid (14.9 g, 53.7 mmol) in THF (500 mL) is added *N,O*-dimethylhydroxylamine hydrochloride (5.24 g, 53.7 mmol), EDC (11.3 g, 59.1 mmol), HOBT (7.98 g, 59.1 mmol) and DIPEA (9.83 mL, 56.4 mmol). The mixture is stirred for about 16 hours at r.t. under nitrogen and then concentrated to
 15 dryness. The resulting residue is taken up in EtOAc, washed with 1M HCl, saturated sodium bicarbonate and brine, and dried (Na₂SO₄). After concentrating to dryness, the resulting residue is purified by flash chromatography (SiO₂, eluting with 1:1 EtOAc/hexane) to give about 12.3g (71%) of the ester. EIS-MS 321 [M+1]

- Step B: 3-Formyl-3,4-dihydro-1H-isoquinoline-2-carboxylic acid tert-
 20 butyl ester
 To a 0°C solution of material from Step A (1.28 g, 4.00 mmol) in THF (30 mL) is slowly added 1.0 M LAH (in THF, 5.1 mL, 5.1 mmol). The reaction mixture is stirred at 0°C for another 15 minutes. To the mixture is slowly added 20 mL of 5% aqueous potassium hydrogensulfate, and the mixture is extracted with Et₂O (2x). The combined organic
 25 portions are washed with 1M HCl, saturated sodium bicarbonate and brine. The mixture is dried (Na₂SO₄) and concentrated to dryness yielding about 0.78g (75%). EIS-MS 262 [M+1]

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5 Step C: 3-{[2-(4-Chloro-phenyl)-1-methoxycarbonyl-ethylamino]-methyl}-
 3,4-dihydro-1H-isoquinoline-2-carboxylic acid tert-butyl ester

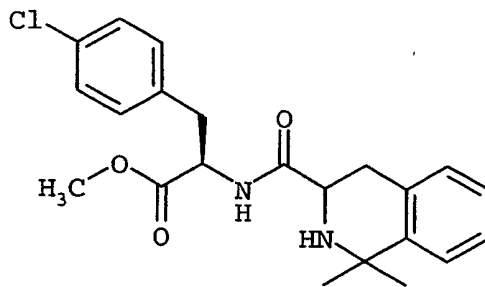
To a 0°C solution of 4-Cl-D-Phe methyl ester (6.27g, 25.1 mmol) and sodium acetate (8.23g, 100.0mmol) in 850ml dry MeOH is added material from Step B (9.8g, 37.6mmol) in 50ml MeOH. The mixture is stirred for about 15 minutes and then sodium
10 cyanoborohydride (2.37g, 37.6 mmol) is added. The cooling bath is removed, and the mixture is stirred for about 16 hours at r.t. The mixture is concentrated to dryness and the resulting residue is taken up in water and 1 mL of 1M HCl. The mixture is extracted with EtOAc and organics are washed with saturated sodium bicarbonate, brine, dried (Na₂SO₄), and concentrated to dryness. The resulting residue is purified by flash
15 chromatography (SiO₂, eluting with 2:1 hexane/EtOAc) to afford about 8.62g (75%).
EIS-MS 459 [M+1]

Step D: To a 12°C solution of material from Step C (1.11g, 2.42mmol) in dioxane (15ml) is added a solution of lithium hydroxide (0.10g, 2.42mmol) in water (7.5 mL). The mixture is stirred for about 16 hours at r.t. and then concentrated to dryness to
20 afford about 1.08g (100%) of the final compound. EIS-MS 445 [M+1]

Preparation 5BC

3-(4-Chloro-phenyl)-2-[(1,1-dimethyl-1,2,3,4-tetrahydro-isoquinoline-3-carbonyl)-
amino]-propionic acid methyl ester

25



To 1,1-dimethyl tic (240 mg, 1.17 mmol) in CH₂Cl₂/DMF (1:1) is added 4-
Cl-D-Phe methyl ester (322 mg, 1.28 mmol), HOBT (197 mg, 1.46 mmol), DIPEA (0.81
mL, 44.68 mmol) and EDC (280 mg, 1.46 mmol). The mixture is stirred at r.t. overnight.
30 The mixture is diluted with EtOAc (100mL) and the organics are washed with saturated

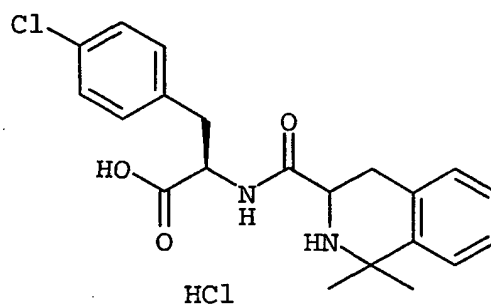
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- 5 aqueous NaHCO_3 , brine, dried (Na_2SO_4), and concentrated to dryness. Purification and separation of the diastereomers is accomplished by flash chromatography (35 g SiO_2 , linear gradient, 40 mL/min 10-50% EtOAc/hexane for 25 minutes and 50% EtOAc/hexane for 7 minutes) to afford about 117 mg of the final compound (25%). LRMS (ESI⁺): 401.1 (M+H).

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Preparation 6BC

3-(4-Chloro-phenyl)-2-[(1,1-dimethyl-1,2,3,4-tetrahydro-isoquinoline-3-carbonyl)-amino]-propionic acid



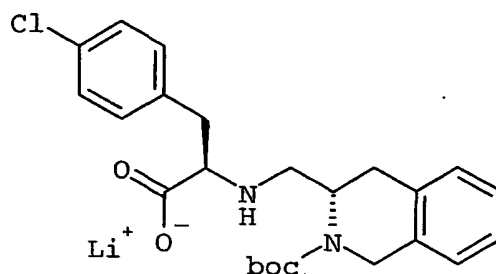
- 15 To the compound 5BC (5.95 g, 14.88 mmol) in a 1:1 mixture of THF/ H_2O (50 mL) is added lithium hydroxide hydrate (0.75 g, 17.87 mmol). The reaction is stirred at r.t. for about 18 hours. The mixture is then concentrated to dryness. The resulting residue is dissolved in water (50 mL) and adjusted to acidic with 1N HCl (25 mL) and then washed with Et_2O (100 mL). The aqueous layer is evaporated to dryness to afford
- 20 about 6.18g (98%) of the final compound. EIS-MS 387 [M+1]

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Preparation 7BC

lithium; 2-[(2-tert-butoxycarbonyl-1,2,3,4-tetrahydro-isoquinolin-3-ylmethyl)-amino]-3-(4-chloro-phenyl)-propionate



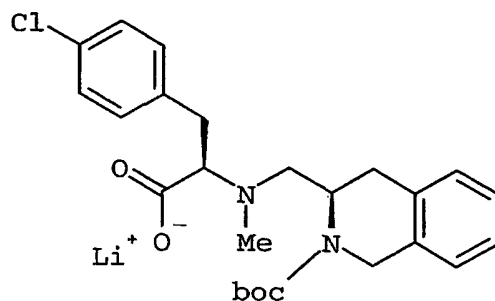
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The above compound was Prepared in a manner similar to the preparation 4BC above except that Boc-L-1,2,3,4-tetrahydroisoquinoline carboxylic acid was used.

Preparation 8BC

15

Preparation of Lithium 2-[(2-tert-butoxycarbonyl-1,2,3,4-tetrahydro-isoquinolin-3-ylmethyl)-methyl-amino]-3-(4-chloro-phenyl)-propionate



20

Step A: To a solution of 3-{[2-(4-Chloro-phenyl)-1-methoxycarbonyl-ethylamino]-methyl}-3,4-dihydro-1H-isoquinoline-2-carboxylic acid tert-butyl ester from preparation 3BC Step C (0.60 gm, 1.31 mmol) in anhydrous methanol, was added sodium acetate (0.54 gm, 6.54 mmol). The solution was brought to pH 5-6 with 3-4 drops of glacial acetic acid. Aqueous formaldehyde (37% by wt., 0.49 mL) was added. The solution was put under a nitrogen atmosphere and cooled to 0°C. After about 15 minutes, sodium cyanoborohydride (0.25 gm, 3.92 mmol) was added and rinsed into the reaction with anhydrous methanol (5 mL). The mixture was stirred at r.t. overnight, and then

25

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- 5 concentrated in vacuo and reconstituted in aqueous sodium bicarbonate and ethyl acetate. After separation of phases, the aqueous phase was extracted with ethyl acetate (2X), and all organics were combined, dried (magnesium sulfate), filtered, and concentrated to an opaque white oil (0.64 gm). Chromatography (0 to 20% ethyl acetate in hexane) gave about 0.6 g of methylated product as a clear oil (97%). MS (m/z, ES⁺): 473.2.
- 10 Step B: A solution of LiOH.H₂O (0.05 gm, 1.27 mmol) in distilled water (4 mL) was added to a solution of the material from Step A in 1,4-dioxane (8 mL), and the reaction was cooled slightly in an ice water bath. The mixture was stirred under a nitrogen atmosphere at r.t. overnight. An additional 1.5 eq. of LiOH.H₂O (0.08 gm) were added as an aqueous solution (4 mL), and the mixture was stirred at r.t. over the weekend.
- 15 The mixture was concentrated, and then combined with THF and concentrated (3x) to help dry the material. The resulting foam was dried at r.t. overnight in a vacuum oven to give about 0.67 g of final compound as a white foam (114%). MS (m/z, ES⁺): 459.2.

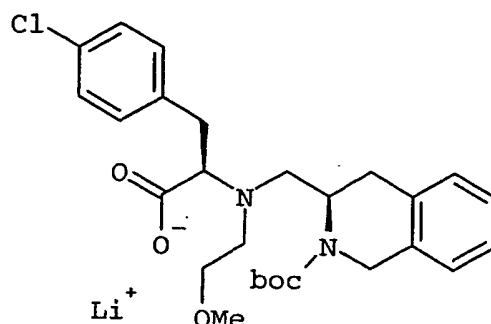
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Preparation 9BC

lithium 2-[(2-tert-butoxycarbonyl-1,2,3,4-tetrahydro-isoquinolin-3-ylmethyl)-(2-methoxy-ethyl)-amino]-3-(4-chloro-phenyl)-propionate



10

Step A: To a solution of methoxyacetaldehyde (0.15 gm, 2.03 mmol), 3-{[2-(4-Chloro-phenyl)-1-methoxycarbonyl-ethylamino]-methyl}-3,4-dihydro-1H-isoquinoline-2-carboxylic acid tert-butyl ester from preparation 3BC Step C (0.31 gm, 0.68 mmol) in acetonitrile was added sodium triacetoxyborohydride (0.72 gm, 3.38 mmol). After stirring overnight under a nitrogen atmosphere at r.t., additional acetaldehyde (0.25 gm) dissolved in acetonitrile and sodium triacetoxyborohydride (0.21 gm) was added, and the mixture was stirred for about 8.5 hours. The mixture was quenched at r.t. with 5N NaOH (5 mL). The aqueous phase was separated from the organic and extracted with ethyl acetate (4x). The combined organics were washed with a brine solution, and then dried, filtered and concentrated. Chromatography (gradient of ethyl acetate in hexane, 0 to 12%) gives about 0.23g of 3-{[2-(4-Chloro-phenyl)-1-methoxycarbonyl-ethyl]- (2-methoxy-ethyl)-amino]-methyl}-3,4-dihydro-1H-isoquinoline-2-carboxylic acid tert-butyl ester as a yellow oil (70%). MS (m/z, ES⁺): 517.2.

25

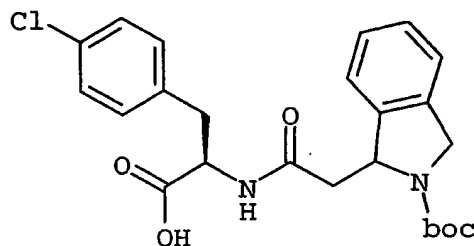
Step B: To a solution of the material from Step A in 1,4-dioxane was added a solution of lithium hydroxide monohydrate (0.05 gm, 1.11 mmol) in distilled water (2 mL). The mixture was stirred overnight at r.t. and then concentrated to a white residue. Addition of THF and concentration (3x) gives the lithium carboxylate as a foam.

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- 5 The foam was dried overnight under vacuum to afford about 0.25g of crude solids (109%). MS (m/z, ES⁺): 503.3.

Preparation 10BC

- 1-[[1-Carboxy-2-(4-chloro-phenyl)-ethylcarbamoyl]-methyl]-1,3-dihydro-isoindole-2-
10 carboxylic acid tert-butyl ester



- Step A: To a suspension of 4-Cl-D-Phe-OMe hydrochloride (40.4 g, 161.5 mmol) in DCM (250 mL) was added saturated aqueous sodium bicarbonate (250 mL),
15 and the mixture was stirred at r.t. for about 1 hour. The organic portion was separated and the aqueous portion was extracted with DCM (2x). The combined organic portions were dried (Na₂SO₄) and concentrated to dryness. To the free amine, in DCM (400 mL) at 0°C, was added 1-Carboxymethyl-1,3-dihydro-isoindole-2-carboxylic acid tert-butyl ester from preparation 2C (isomer 2, 44.8 g, 161.5 mmol), EDC (31.0 g, 161.5 mmol) and
20 4-DMAP (2.0 g, 16.1 mmol). The mixture was stirred at 0°C for about 30 minutes whereupon the cooling bath was removed and the mixture was stirred for another 5 hours at r.t. The mixture was then washed with saturated aqueous sodium bicarbonate (200 mL) and 10% aqueous sodium bisulfate (200 mL), and then dried (Na₂SO₄) and concentrated to dryness to afford about 76.4 g (100%) of the ester. EIS-MS 471 [M-1].

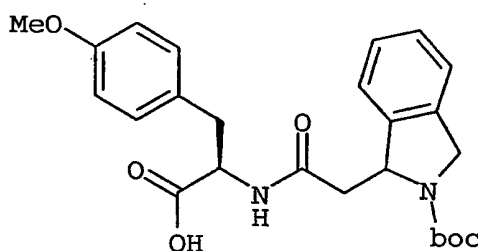
- 25 Step B: To the ester from Step A (76.4 g, 161.5 mmol) in MeOH (760 mL) was added 1 N NaOH (242.0 mL, 242.0 mmol), and the mixture was heated at 50°C for 4 hours and then stirred for another 16 hours at r.t. After concentrating to dryness, the resulting residue was taken up in 500 mL of water and washed with diethyl ether (2x). The aqueous portion was acidified to pH 2 with 10% aqueous sodium bisulfate and
30 extracted with EtOAc (4 x 200 mL). The combined organic extracts were dried (MgSO₄)

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- 5 and concentrated to dryness. The resulting solid was suspended in hexanes, filtered, and dried to afford about 67.7 g (91%) of the final compound. EIS-MS: 457 [M-1].

Preparation 11BC

- 10 1-{[1-Carboxy-2-(4-methoxy-phenyl)-ethylcarbamoyl]-methyl}-1,3-dihydro-isoindole-2-carboxylic acid tert-butyl ester



- 15 Step 1: To a solution of p-methoxy-D-Phe-OMe (1.72g, 8.23 mmol) dissolved in THF (45mL) and 1-carboxymethyl-1,3-dihydro-isoindole-2-carboxylic acid tert-butyl ester (2.51g, 9.05mmol) was added HOBT (1.22g, 9.05mmol), EDC (1.73g, 9.05mmol) and DIPEA (1.6mL, 9.05mmol). The reaction was stirred overnight at r.t. and then concentrated. The mixture was washed with 1M HCl, dilute NaHCO₃ and brine, and then
- 20 dried with sodium sulfate. The mixture was chromatographed on silica gel eluting with 3% 2M NH₃ in MeOH/CH₂Cl₂ giving about 2.58g as white solids. Mass MH⁺ 469

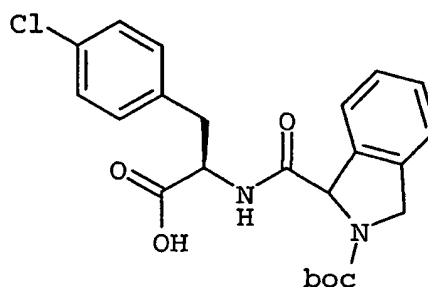
- Step 2: The white solid from Step 1 (2.58g, 5.5 mmol) was dissolved in dioxane (37 mL) and lithium hydroxide hydrate (0.35g, 8.3mmol) dissolved in H₂O (19 mL) was added. The mixture was stirred for about 2.5 hours at r.t. and then concentrated.
- 25 Ethyl acetate was added and the mixture was treated with 1M HCl, which was then washed with brine and concentrated to afford about 2.56 g of the final free acid. LRMS (ESI⁺): 455 (M+1)

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Preparation 12BC

1-[1-Carboxy-2-(4-chloro-phenyl)-ethylcarbamoyl]-1,3-dihydro-isoindole-2-carboxylic acid tert-butyl ester



10

Step 1: About 2.0g (7.60 mmol) of (R,S)-Boc-1,3-dihydro-2H isoindole carboxylic acid was dissolved in 100ml THF and about 2.28 g (9.12mmol) of 4-Cl-D-phe-methylester HCl, 1.25g (9.12mmol) of HOBT, 1.75g (9.12mmol) of EDC, and 1.6ml (9.12mmol) of DIEA were added. The mixture was stirred overnight at r.t., concentrated to dryness, washed with 1M HCl, dilute NaHCO₃ and brine, and then dried over sodium sulfate. The material was chromatographed on silica gel by eluting with ethyl acetate/hexane 1:2 to give about 1.05g of isomer 1 and about 0.82g of isomer 2, and about 1.61g mixture of isomers 1 and 2. Mass MH⁺ 459

15

20

Step 2: About 0.82g (1.79 mmol) of the isomer 2 obtained in Step 1 was dissolved in 11ml of dioxane and 0.11g (2.68 mmole) of LiOH-hydrate in 5.5 ml of H₂O was added. The mixture was stirred for about 4 hours at r.t. and then concentrated to dryness. Ethyl acetate was added, and the solution was washed with 1M HCl and brine, and then concentrated to dryness affording about 0.75 g of the free acid. Mass: 445 (MH⁺).

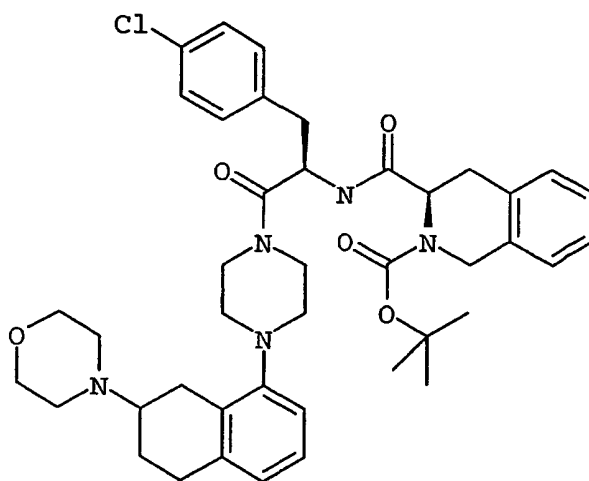
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5 EXAMPLES

Example 1

3-(4-chloro-phenyl)-2-[(1,1-dimethyl-1,2,3,4-tetrahydroisoquinoline-3-carbonyl)-amino]-
propionic acid



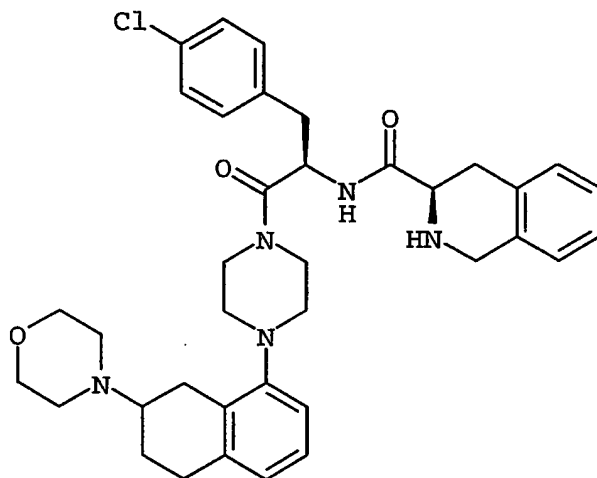
10 To compound 4A (0.441g, 0.96mmol), in DCM (8mL) and DMF (2mL), is
added compound 1BC (0.289g, 0.96mmol), DIPEA (1.6mL, 9.60mmol), and HATU
(0.365g, 0.96mmol). The mixture is stirred at r.t. overnight and then concentrated to
dryness. The resulting residue is taken up in EtOAc (50mL), and the organics are washed
15 with saturated NaHCO₃, H₂O and brine, and then concentrated to dryness. The desired
product is purified by flash chromatography (SiO₂, eluting with EtOAc-TEA-MeOH,
98:1:1 to 90:5:5) to afford about 0.60g of the final compound (84%). EIS-MS 742.3
[M+1]

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Example 2

1,2,3,4-Tetrahydro-isoquinoline-3-carboxylic acid {1-(4-chloro-benzyl)-2-[4-(7-morpholin-4-yl-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-2-oxo-ethyl}-amide



To compound of Example 1 (0.596g, 0.803mmol), in DCM (5mL), is
10 added TFA (5mL) and the mixture is stirred at room temperature for about 5 minutes.
After concentrating to dryness, the resulting residue is triturated with Et₂O, and the
resulting solid is collected by filtration and dried to afford about 0.52g of the final
compound (91%). EIS-MS 642.2 [M+1]

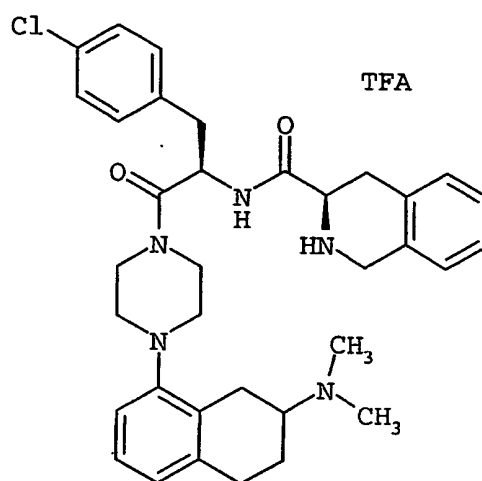
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Example 3

1,2,3,4-Tetrahydro-isoquinoline-3-carboxylic acid {1-(4-chloro-benzyl)-2-[4-(7-dimethylamino-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-2-oxo-ethyl}-amide;
trifluoroacetate



10

Step 1: To the A domain compound described in Preparation 1A (917mg, 3.75mmol) is added N-Boc-D-Tic-4-Cl-D-Phe-OH (1.7g, 3.75mmol), HATU (1.43g, 3.75mmol), DIPEA (1.3mL, 7.5mmol), CH₂Cl₂ (10mL) and DMF (3mL). The mixture is stirred at r.t. for 12 hours and then diluted with EtOAc (15 fold). The organics are washed with saturated NaHCO₃, H₂O and brine, and then concentrated to dryness. The resulting residue is purified by flash chromatography (Silica gel 60, eluting with EtOAc) and then used in Step 2 below (917mg, 35%). Ion spray MS: 770.2 [M+].

15

Step 2: To the compound of Step 1 (917mg, 1.30mmol) in CH₂Cl₂ (10mL) is added TFA (8mL), and the mixture is stirred at r.t. for about 30 minutes. After concentrating to dryness, the resulting residue is triturated with Et₂O, and the TFA salt is collected via filtration and dried under vacuum to afford about 920 mg of the final compound (85%). Ion spray MS: 600.2 [M+].

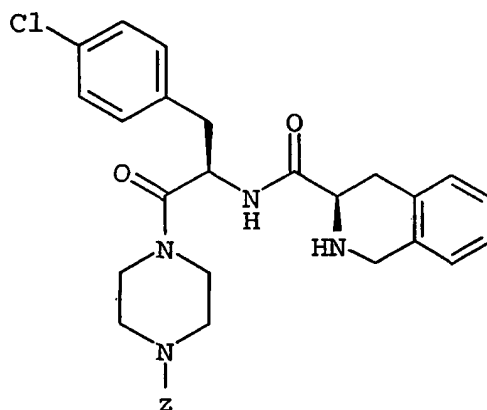
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Examples 4-18

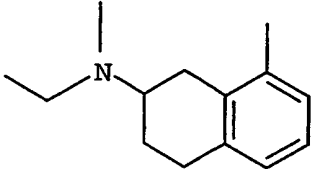
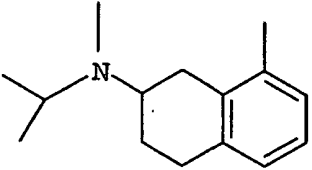
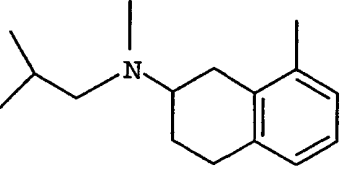
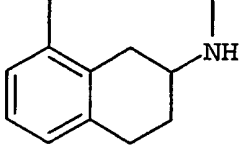
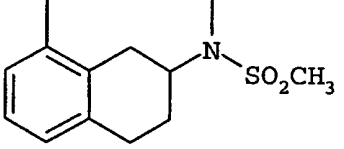
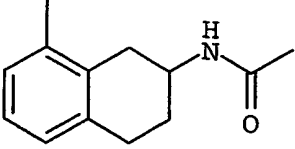
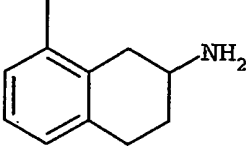
The compounds of examples 4-18 are prepared from using appropriate A domain and the compound 1BC (Preparation 1BC) by following a substantially similar procedure as described in Examples 1 and 2.



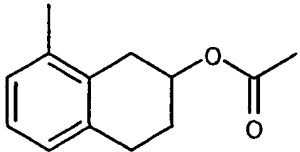
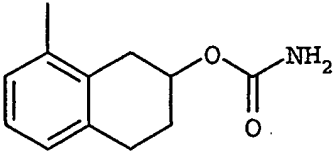
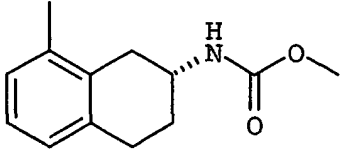
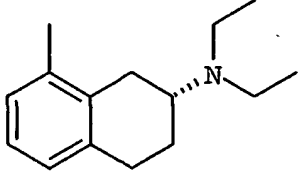
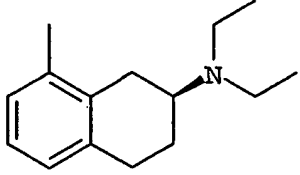
10

No	Z	EIS-MS [M+1]
4		625.2
5		656.2
6		626.2

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No	Z	EIS-MS [M+1]
7		614.2
8		628.2
9		642.3
10		586.1
11		664.2
12		614.1
13		572.2

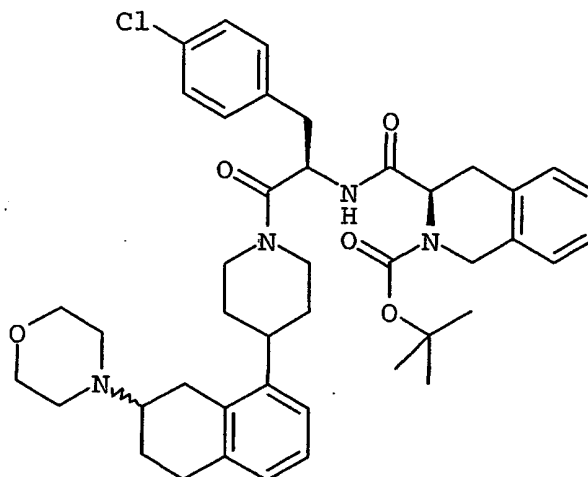
-163-

No	Z	EIS-MS [M+1]
14		615.2
15		615.2
16		630.2
17		629.1
18		629.1

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Example 19

3-{1-(4-chloro-phenyl)-2-[4-(7-morpholin-4-yl-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperidin-1-yl]-2-oxo-ethylcarbamoyl}-3,4-dihydro-1H-isoquinolin-2-carboxylic acid
tert-butyl ester



To the compound 34A (0.351g, 0.779mmol), in DCM (8mL) and DMF (2mL), is added compound 1BC (0.234g, 0.779mmol), DIPEA (1.36mL, 7.79mmol), and HATU (0.296g, 0.779mmol). The mixture is stirred at r.t. overnight and then concentrated to dryness. The resulting residue is taken up in EtOAc (50mL) and the organics washed with saturated NaHCO₃, H₂O, brine, and concentrated to dryness. The desired product is purified by flash chromatography (SiO₂, eluting with 100% EtOAc to EtOAc-TEA-MeOH, 90:5:5) to afford about 0.46g of the final product (79%). EIS-MS 741.3 [M+1]

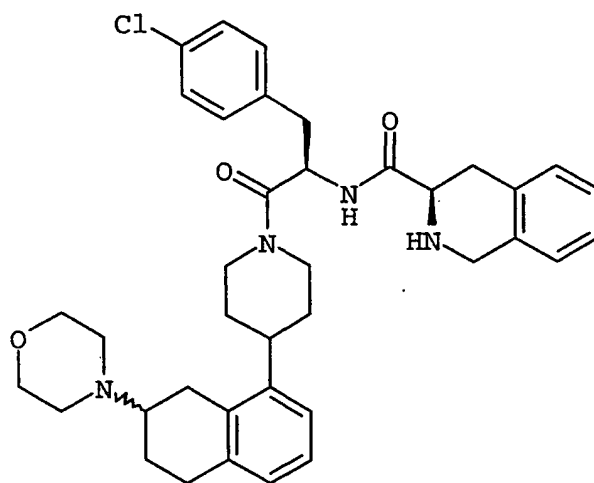
20

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5

Example 20

1,2,3,4-Tetrahydro-isoquinoline-3-carboxylic acid {1-(4-chloro-benzyl)-2-[4-(7-
morpholin-4-yl-5,6,7,
8-tetrahydro-naphthalen-1-yl)-piperidin-1-yl]-2-oxo-ethyl}-amide



10

To compound prepared in Example 19 (0.42g, 0.567mmol), in DCM (3mL), is added TFA (3mL) and the mixture is stirred at room temperature for about 45 minutes. After concentrating to dryness, the resulting residue is triturated with Et₂O, and the resulting solid is collected by filtration and dried to afford about 0.47g of the final product (94%). EIS-MS 641.2 [M+1]

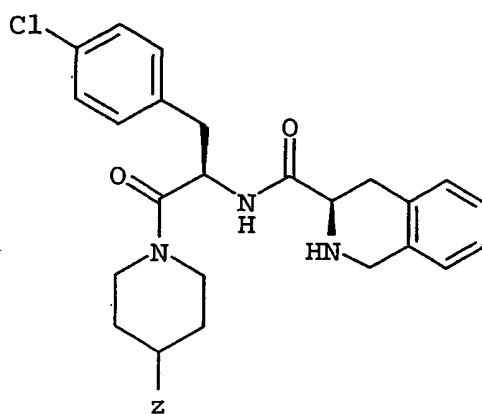
15

-166-

5

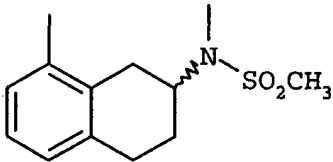
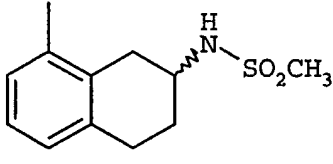
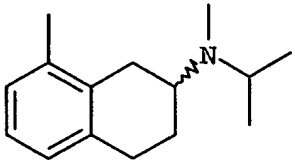
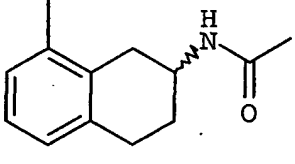
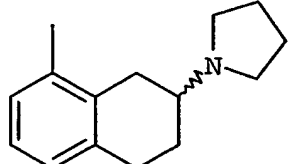
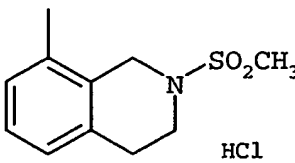
Examples 21-29

The compounds of examples 21-29 are prepared from the appropriate A domain and the compound 1BC by following a substantially similar procedure as described in Examples 19 and 20.



No	Z	(EIS)-MS [M+1]
21		627.3
22		655.3
23		599.2

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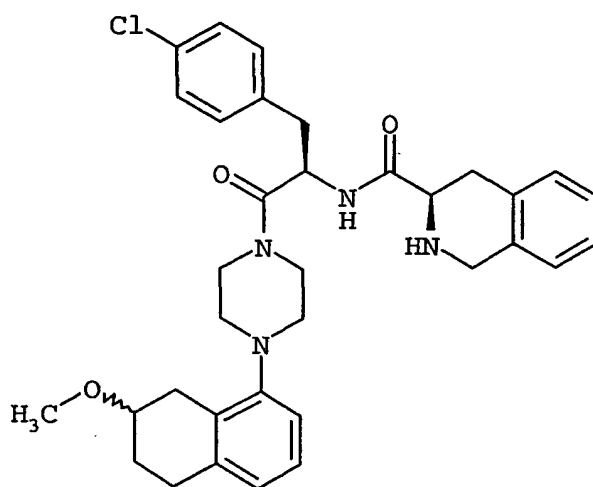
No	Z	(EIS)-MS [M+1]
24		664.2
25		649.1
26		627.3
27		613.1
28		625.2
29		635.1

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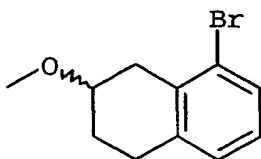
5

Example 30

1,2,3,4,-tetrahydro-isoquinoline-3-carboxylic acid{1-(4-chlorobenzyl)-2-[4-(7-methoxy-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-2-oxo-ethyl}amide



10

Step 1:

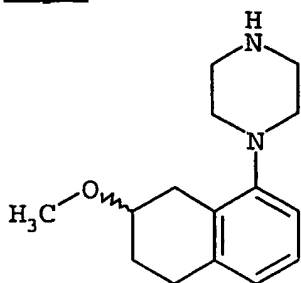
15

To 8-Bromo-2-hydroxytetralin (1.07g, 4.71mmol), prepared from 8-bromo-2-tetralone by essentially following the procedure outlined in *J. Org. Chem.*, 56(19): 5564-5566, 1991., in THF (5mL), is added NaH (60% dispersion in oil, 330mg, 8.24mmol) and the mixture is stirred at r.t. for about 30 minutes. Iodomethane (367 microliters, 5.89 mmol) was then added and after about an hour the mixture is quenched by the cautious addition of H₂O (10mL) and the desired product is extracted into EtOAc (50mL). The organic extracts are concentrated to dryness and the desired methyl ether is purified by flash chromatography (SiO₂, eluting with toluene) to afford about 0.94g of the final product (83%). EIS-MS 242.1 [M+1]

20

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5

Step 2:

The A domain compound is prepared from the compound of Step 1 and N-Boc-piperazine by following substantially similar procedure as described in Preparation 3A and 4A. EIS-MS 247.2 [M+1]

10

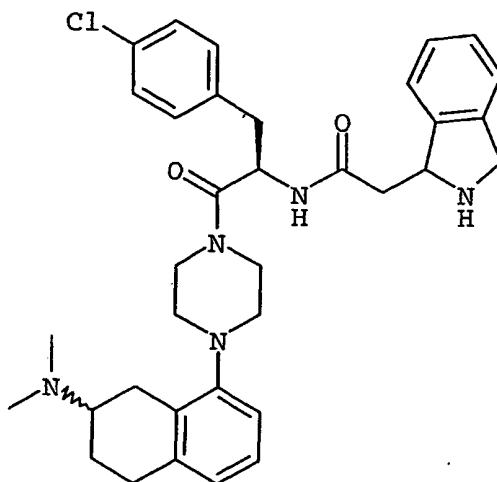
The final compound of Example 31 is prepared by coupling the compound of step 2 and the compound 1BC by following substantially similar procedure as described in Examples 19 and 20. EIS-MS 587 [M+1]

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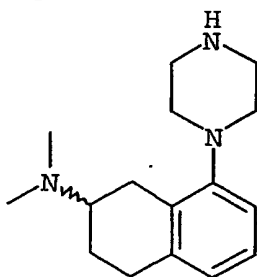
5

Example 31

N-{1-(4-chloro-benzyl)-2-[4-(7-dimethylamino-5,6,7,8-tetrahydro-naphthalen-1-yl)-
piperazin-1-yl]-2
-oxo-ethyl}-(2,3-dihydro-1H-isoindol-1-yl)-acetamide



10

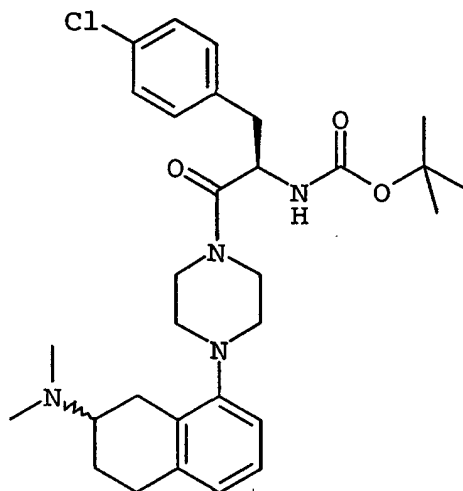
Step 1:

The Boc-protected title compound is prepared from 2-dimethylamino-8-bromo-1,2,3,4-tetrahydronaphthalene (European Patent Application 91-304047, 1.0g, 3.93 mmol) and commercially available 1-Boc-piperazine (0.805g, 4.32 mmol) by following the conditions outlined in *J. Am. Chem. Soc.*, 118:7215-7216, 1996. The desired product is purified by flash chromatography (SiO₂, eluting with 98:1:1, EtOAc-TEA-MeOH) to afford about 1.35g (95%). EIS-MS 360.2 [M+1] To the Boc-protected compound (1.35g, 3.75 mmol) is added TFA (5mL) and DCM (5mL) and the resulting mixture is stirred at r.t. for about 30 minutes. After concentrating to dryness the resulting residue is taken up in 1N NaOH and the desired amine is extracted into EtOAc (100 mL).

-171-

- 5 The organics are washed with H₂O, brine, and re-concentrated to afford about 970 mg (quantitative). EIS-MS 260.1 [M+1]

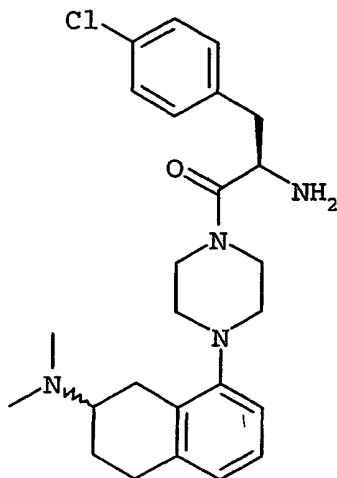
Step 2:



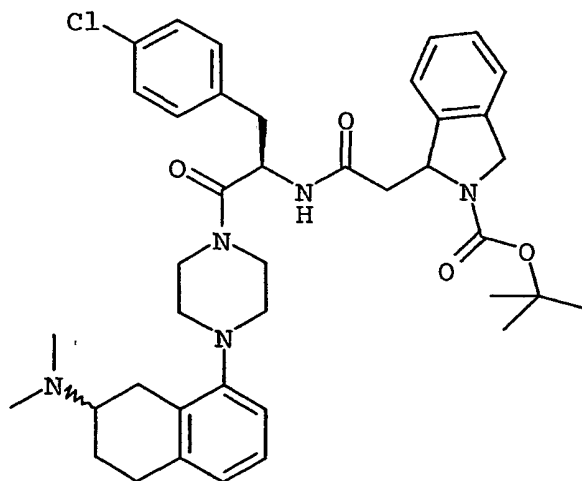
- 10 To the compound of Step 1 (0.68 g, 2.62 mmol) is added N-Boc-4-Cl-D-Phe (0.785g, 2.62 mmol), EDC (0.553g, 2.88 mmol), 4-DMAP (catalytic), DCM (4mL), and DMF (1mL). The mixture is stirred at r.t. for about one hour and then diluted with EtOAc (50mL). The organics are washed with saturated NaHCO₃, H₂O, brine and concentrated to dryness. The resulting residue is purified by flash chromatography (SiO₂,
15 eluting with EtOAc-TEA-MeOH, 90:5:5). EIS-MS 541.2 [M+1]

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5

Step 3:

To the compound of Step 2 (1.41g, 2.62 mmol), in DCM (5mL), is added TFA (5mL) and the mixture is stirred at r.t. for about 45 minutes. After concentrating to dryness, the resulting residue is taken up with 1N NaOH and the desired "free" amine is extracted into EtOAc (100mL). The organic extracts are concentrated to dryness to afford about 1.1g (96%). EIS-MS 441.2 [M+1]

Step 4:

15

To the compound of Step 3 (0.279 g, 0.632 mmol) is added compound 5C (isomer 2, 0.175g, 0.632 mmol), EDC (0.133g, 0.695 mmol), 4-DMAP (catalytic), and DCM (6mL). The mixture is stirred at r.t. for about an hour and then diluted with EtOAc (50 mL). The organics are washed with saturated NaHCO₃, H₂O, brine, and concentrated

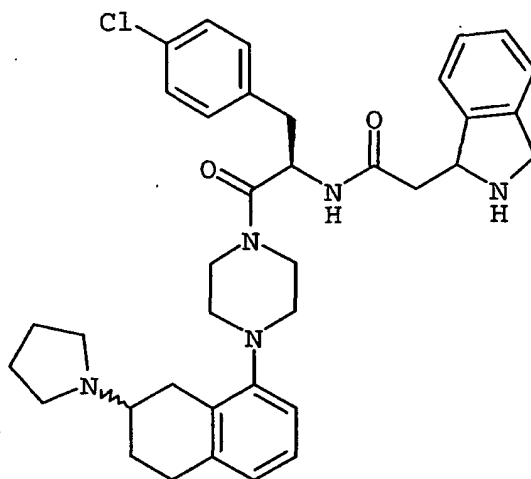
-173-

5 to dryness. The resulting residue is purified by flash chromatography (SiO₂, eluting with EtOAc-TEA-MeOH, 90:5:5) to afford 430 mg (97%). EIS-MS 700.2 [M+1]

Step 5: To the compound of Step 4 (0.430 g, 0.614 mmol), in DCM (5mL), is added TFA (5 mL) and the mixture is stirred at r.t. for about 45 minutes. After concentrating to dryness, the resulting residue is triturated with Et₂O, and the resulting
10 solid is collected by filtration and dried to afford about 0.325 g of the final product (74%). EIS-MS 600.2 [M+1]

Example 32

N-{1-(4-Chloro-benzyl)-2-oxo-2-[4-(7-pyrrolidin-1-yl-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-ethyl}-2-(2,3-dihydro-1H-isoindol-1-yl)-acetamide
15



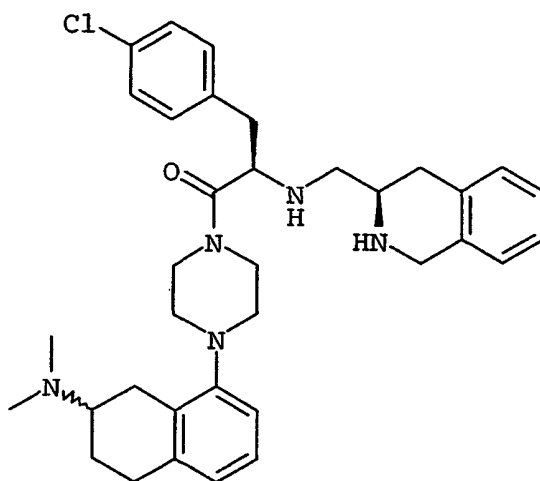
Example 32 is prepared from the compound 8A by following a substantially similar procedure as described in Example 31. EIS-MS 624.4 [M-1].

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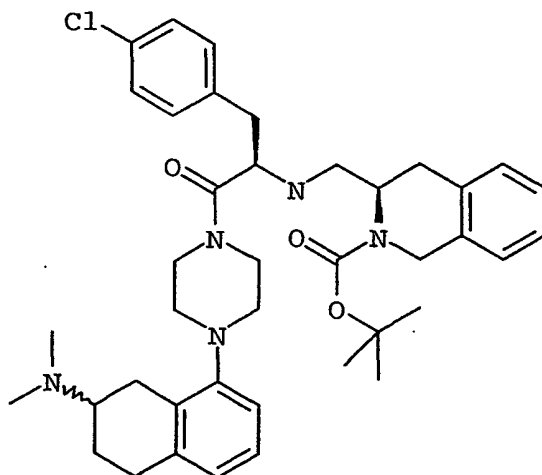
5

Example 33

3-(4-Chloro-phenyl)-1-[4-(7-dimethylamino-5,6,7,8-tetrahydro-naphthalen-1-yl)-
piperazin-1-yl]-2-[1,
2,3,4-tetrahydro-isoquinolin-3-ylmethyl)-amino]-propan-1-one



10

Step 1:

To the compound obtained in Step 1 of Example 31 (0.150g, 0.685 mmol),
15 in DCM, is added compound 4BC (0.307g, 0.685 mmol), EDC (0.131 g, 0.685 mmol) and
HOBT (0.106 g, 0.685 mmol). The mixture is stirred at r.t. for about 4 hours and then
diluted with EtOAc (50 mL). The organics are washed with saturated NaHCO₃, H₂O,
brine, and concentrated to dryness. The resulting residue is purified by flash

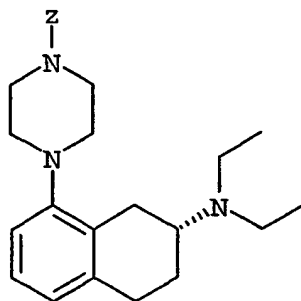
-175-

- 5 chromatography (SiO₂, eluting with 5% 2.0 M ammonia in MeOH-DCM) to afford about 210 mg (45%). EIS-MS 686 [M+1]

Step 2: To the compound of Step 1 (0.20 g, 0.29 mmol), in CH₂Cl₂ (5 mL), is added TFA (5 mL) and the mixture is stirred at r.t. for about 45 minutes. After concentrating to dryness, the resulting residue is triturated with Et₂O, and the resulting
 10 solid is collected by filtration and dried to afford about 0.26g of the final product (97%).
 EIS-MS 586.3 [M+1]

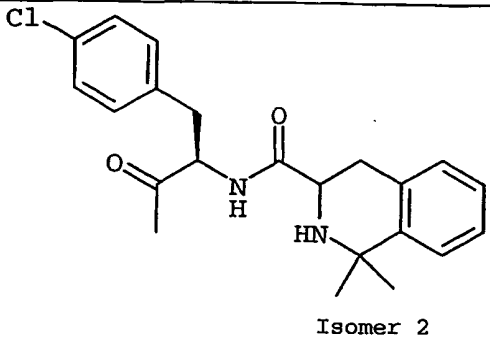
Examples 34-35 are prepared from the compound 30A and the appropriate B-C domain by following the substantially similar procedures described in Examples 1 and 2.

15

Examples 34-35

No	Z	EIS-MS [M+1]	BC Domain
34	<p>Isomer 1</p>	656.3	6BC Isomer 1

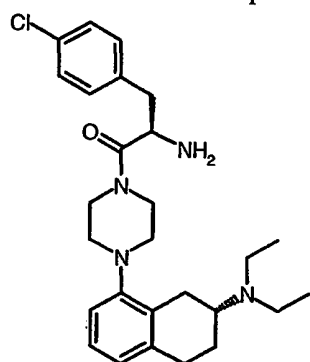
-176-

No	Z	EIS-MS [M+1]	BC Domain
35	 Isomer 2	656.3	6BC Isomer 2

5

Examples 36-37

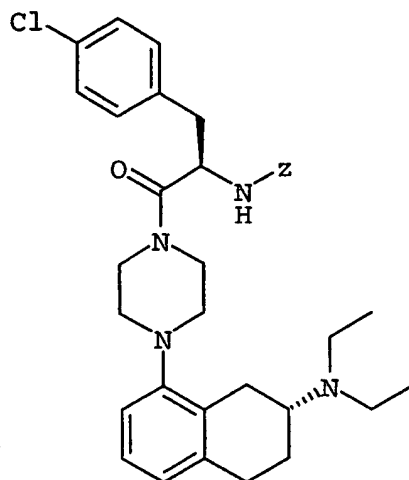
Examples 36 and 37 are prepared from the amine compound,



10

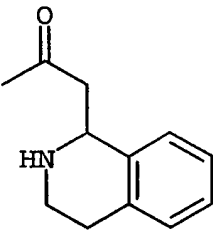
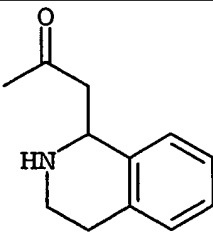
, which is prepared from the compound 30A and 4-Cl-D-Phe by following a substantially similar procedure as described in Example 31, Steps 2 and 3 (62%). EIS-MS 569.1 [M+1]

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5

and the appropriate B-C domain by following a substantially similar procedure as described in Example 31, Steps 4 and 5.

No	Z	EIS-MS [M+1]	C Domain
36	 Isomer 1	642.1	6C Isomer 1
37	 Isomer 2	642.1	6C Isomer 2

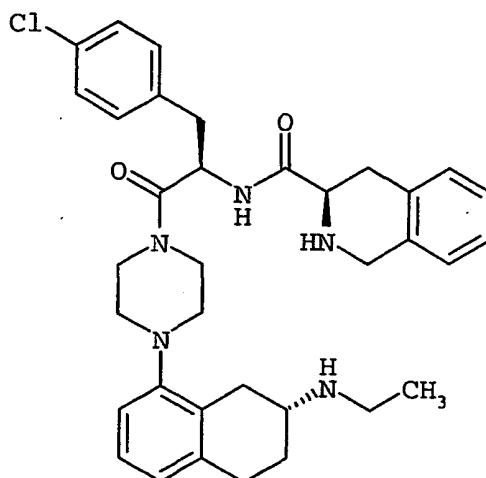
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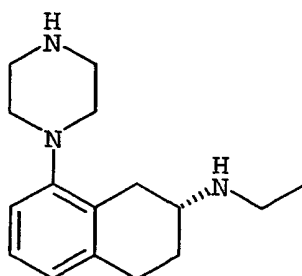
5

Example 38

1,2,3,4-tetrahydro-isoquinoline-3-carboxylic acid{1-(4-chlorobenzyl)-2-[4-(7-ethylamino-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-2-oxo-ethyl}amide



10

Step 1:

To the compound 26A (4.26g, 12.85 mmol), in DMF (30 mL), is added ethyl bromide (2.89 mL, 38.55 mmol). The mixture is stirred at r.t. for about 48 hours, and an additional equivalent of ethyl bromide is added. The mixture is stirred for another 24 hours. The potassium carbonate is removed by filtration and washed with EtOAc. The filtrate is washed with H₂O, and the organics are dried over anhydrous Na₂SO₄ and concentrated to dryness. The resulting residue is purified by flash chromatography (SiO₂, eluting with EtOAc-TEA-MeOH, 98:1:1) to afford about 2.3g (46%) diethyl amine and about 0.183g (4%) mono-ethyl amine.

20

Step 2: The mono-ethyl amine is taken up in DCM (5 mL) and TFA (5 mL) is added. The mixture is stirred at r.t. for about 45 minutes. After concentrating to

-179-

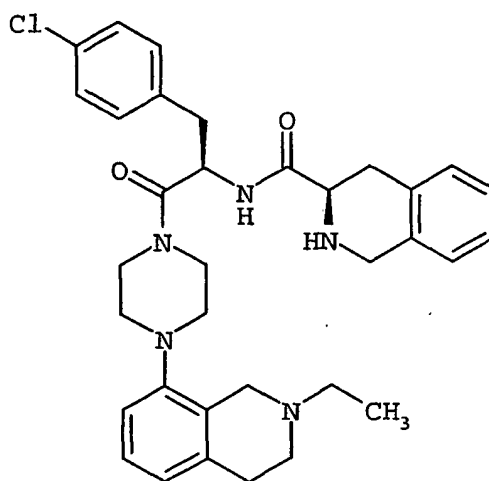
5 dryness, the resulting residue is taken up with 1N NaOH and the desired "free" amine is extracted into EtOAc (50 mL). The organic extracts are concentrated to dryness.

Step 3: The final compound is prepared by coupling the compound obtained above and the compound 1BC by following a substantially similar procedure as described in Examples 1 and 2 to afford about 181 mg (53%). EIS-MS 600.1 [M+H].

10

Example 39

1,2,3,4,-tetrahydro-isoquinoline-3-carboxylic acid{1-(4-chlorobenzyl)-2-[4-(2-ethyl-1,2,3,4-tetrahydro-isoquinolin-8-yl)-piperazin-1-yl]-2-oxo-ethyl}amide



15

To compound 48A (226 mg, 0.92 mmol) and the compound 1BC (423 mg, 0.92 mmol), in DCM (10 mL), are added HATU (350mg, 0.92 mmol) and DIPEA (0.323 mL, 1.84 mmol). The reaction is stirred at r.t. for about 12 hours and then concentrated to dryness. The resulting residue is purified by flash chromatography on silica gel. The coupled product is dissolved in a 1:1 mixture of DCM and TFA (10 mL) and stirred at r.t. for about 2 hours. After concentrating to dryness, the resulting residue is subjected to SCX ion exchange chromatography followed by flash chromatography on silica gel. The resultant foam is dissolved in DCM (10 mL) and 2.0 M HCl in diethyl ether added (5 mL). The slurry is concentrated to dryness, and the resultant solid dried under vacuum for 12 hours to give afford about 260mg of the final compound (43%). EIS-MS 586.3 [M+1]

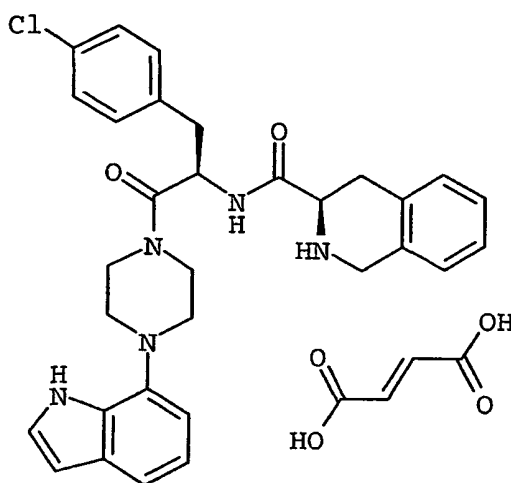
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5

Example 40

1,2,3,4,-tetrahydro-isoquinoline-3-carboxylic acid {1-(4-chloro-benzyl)-2-[4-{1H-indol-7-yl}-piperazin-1-yl]-2-oxo-ethyl}-amide



10 N-Boc-D-Tic-4-Cl-D-Phe-OH (100mg, 0.22mmol), the A domain compound of Preparation 69A (45mg, 0.22mmol) and HATU (84mg, 0.22mmol) are dissolved in 1mL of 10% DMF in DCM followed by the addition of DIPEA (77 microliters, 0.44mmol). The mixture is stirred under N₂ at r.t. for about 2 hours and then concentrated under vacuum. The residue is taken up in DCM (2mL) and TFA is (2mL)

15 added. The mixture is stirred for about 2 hours and then concentrated to an oily solid. The mixture is "free-based" via SCX resin column and then chromatographed on silica gel. Product containing fractions are combined to give "free-based" material (46mg). Fumaric acid (about 1 eq.) in methanol (10mL) is added and the mixture is concentrated, triturated in ether, filtered and dried to give about 23 mg (16%) of the final product as the

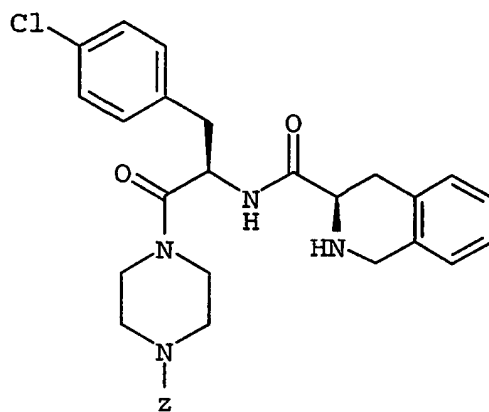
20 fumarate salt. EIS-MS: Found 542.3 M+1.

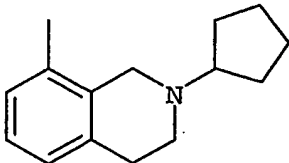
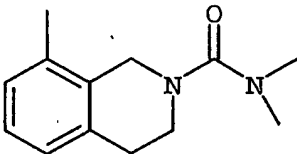
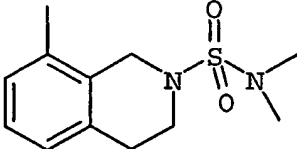
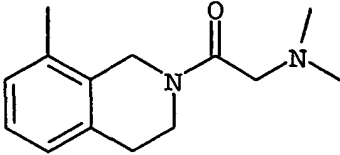
-181-

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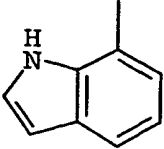
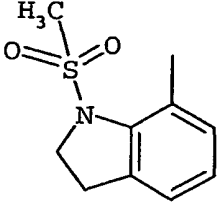
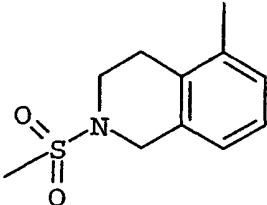
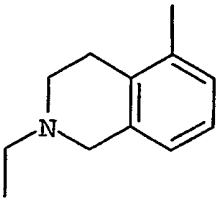
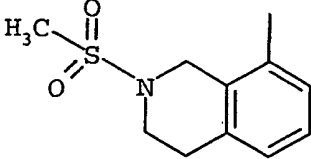
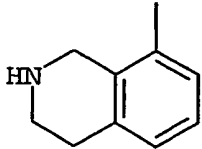
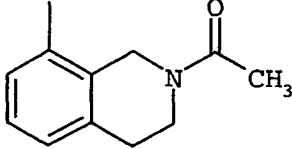
Examples 41-64

Examples 41-64 are prepared from the appropriate A domains and the compound 1BC by following a substantially similar procedure as described in Example 30.

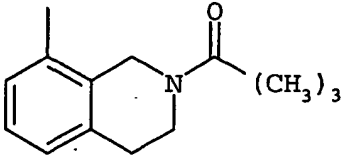
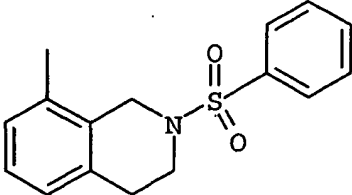
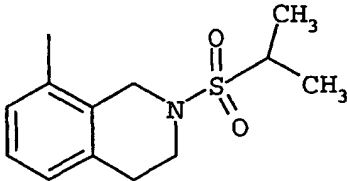
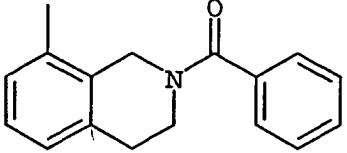
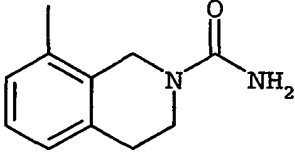
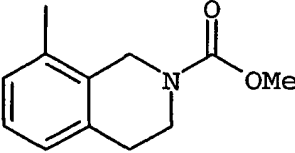
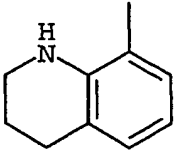


No	Z	EIS-MS [M+1]
41		626.3
42		629.3
43		665.2
44		643.3

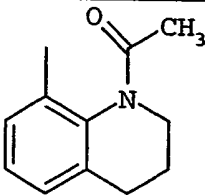
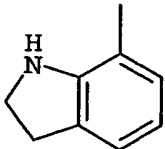
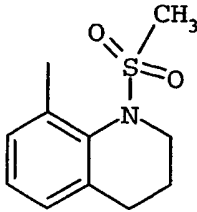
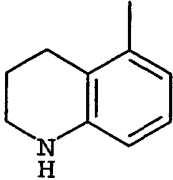
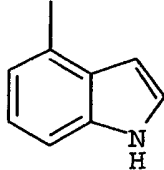
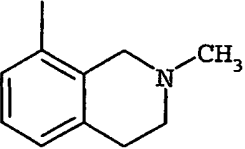
-182-

No	Z	EIS-MS [M+1]
45	 <chem>Cc1ccc2c(c1)c[nH]2</chem>	542.3
46	 <chem>Cc1ccc2c(c1)CN(C2)S(=O)(=O)C</chem>	622.3
47	 <chem>Cc1ccc2c(c1)CN(C2)S(=O)(=O)C</chem>	636.3
48	 <chem>Cc1ccc2c(c1)CN(C2)CC</chem>	586.3
49	 <chem>Cc1ccc2c(c1)CN(C2)S(=O)(=O)C</chem>	636.1
50	 <chem>Cc1ccc2c(c1)CN(C2)CC</chem>	558.5
51	 <chem>Cc1ccc2c(c1)CN(C2)CC(=O)C</chem>	600.2

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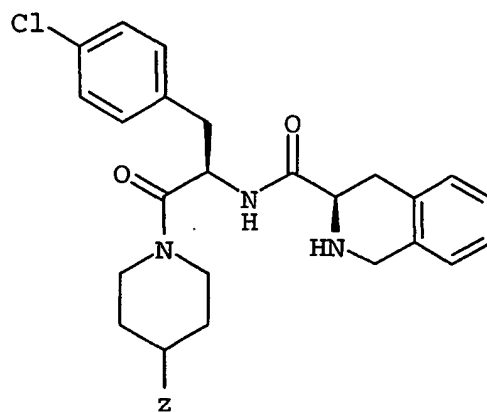
No	Z	EIS-MS [M+1]
52		642.2
53		698.0
54		664.1
55		662.1
56		601.2
57		616.3
58		558.3

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No	Z	EIS-MS [M+1]
59	 <chem>CC(=O)N1CCc2ccccc2N1</chem>	600.2
60	 <chem>CC1CN2C=CC=CC=C2CC1</chem>	544.3
61	 <chem>CC(=O)N1CCc2ccccc2N1S(=O)(=O)C</chem>	636.1
62	 <chem>CC1CN2C=CC=CC=C2CC1</chem>	558.3
63	 <chem>CC1CN2C=CC=CC=C2CC1</chem>	542.3
64	 <chem>CCN1CCc2ccccc2CC1</chem>	572.1

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- 5 Examples 65-68 are prepared from the appropriate A domains and the compound 1BC by following a substantially similar procedure as described in Example 39.



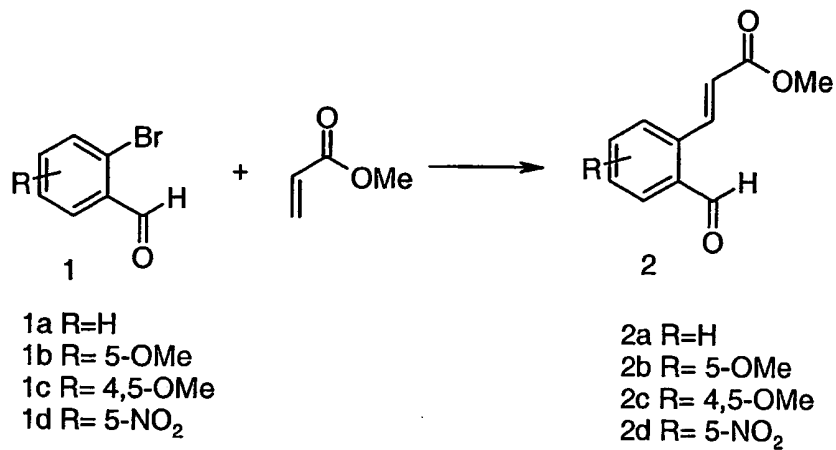
No	Z	EIS-MS [M+1]
65		599.1
66		585.2
67		571.0
68		635.2

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Preparation of Novel C-Domain Pieces

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5 Heck Coupling:



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Preparation PP1

Synthesis of Compound (2a) by a Heck Coupling of 2-bromobenzaldehyde (1a) with methyl acrylate ($\text{Pd}(\text{OAc})_2/\text{PPh}_3$ as the catalyst): A mixture of 2-bromobenzaldehyde (1a) (24.5 g, 132 mmol), methyl acrylate (17.9 mL, 199 mmol), $\text{Pd}(\text{OAc})_2$ (590 mg, 2.65 mmol, 2 mol%), PPh_3 (1.39 g, 5.30 mmol, 4 mol%) and Et_3N (46 mL, 331 mmol) was stirred at 80 °C for 15 h. Large amount of yellow solid was formed after the reaction was done. The mixture was cooled to r.t., concentrated, and mixed with H_2O (200 mL). The organic solid was collected by filtration, and then applied to a plug of silica gel (25 g) (EtOAc /hexane 1:1) to give a dark yellow solid. The solid was purified by crystallization (100 mL EtOAc bottom layer, 120 mL hexane top layer) to provide 17.57 g (70%) (100 % pure by NMR) of the first crop and 5.23 g (21%) (95% by NMR) of the second crop of 2a.

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Preparation PP2

Synthesis of Compound (2a) by a Heck Coupling of 2-bromobenzaldehyde (1a) with Methyl Acrylate ($\text{R}=\text{H}$) ($\text{Pd}(\text{OAc})_2/\text{P}(\text{O-Tolyl})_3$ as the catalyst): The compound 1a (9.998 g, 54.04 mmol) was dissolved in toluene (20 mL) at r.t. Methylacrylate (5.996 g, 69.65 mmol, 1.29 eq.), NEt_3 (15 mL), $\text{Pd}(\text{OAc})_2$ and $\text{P}(\text{O-Tolyl})_3$ were successively added and the mixture was stirred under reflux. After 2 hours, the reaction mixture was allowed to cool to r.t. Then the precipitated yellow catalyst was removed by filtration. The catalyst was rinsed with toluene (2 X 10 mL) and the filtrates were concentrated to dryness under reduced pressure. The residual oil was dried under vacuum over the weekend to give a crude solid (11.449 g). The solid was taken-up with isopropanol (25 mL) and stirred overnight at r.t. Then, the precipitate was filtered and rinsed with isopropanol (5 mL). The wet cake (8.240 g) was dried overnight at r.t. affording the highly pure 2-carboxaldehyde-methyl-cinnamate with 74% yield (7.627 g, 40.1 mmol).

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Preparation PP3

Heck Coupling of 1b and methyl acrylate to form 2b (R = 5-OMe): A mixture of 2-bromo-5-methoxybenzaldehyde (1b) (4.5 g, 20.9 mmol, Aldrich), methyl acrylate (2.7 g, 1.5 eq, 2.83 mL), Et₃N (7.4 g, 3.5 eq, 10.2 mL), Pd(OAc)₂ (93 mg, 0.02 eq), and P(O-Tol)₃ was stirred and heated to 80°C over 2-3 days. The reaction mixture was cooled to r.t., partitioned between EtOAc (50 mL) and brine (50 mL). The aqueous was extracted with EtOAc (2x50 mL). The combined organic was washed with brine (1x50 mL), dried over MgSO₄, filtered, concentrated to yield a yellow brown oil (5.01 g, 109 %). This crude oil was purified in a hot solvent Hex / EtOAc (80 mL/ 15 mL) to yield 2b as a pale yellow solid (3.5 g, 76 %).

15

Preparation PP4

Heck Coupling of 1c and Methyl Acrylate to Form 2c (R = 4,5-OMe): To a solution of 1c (906 mg, 3.70 mmol) in toluene (2 mL) was added Pd(OAc)₂ (17 mg, 0.074 mmol, 2 mol%), P(O-Tolyl)₃ (45 mg, 0.148 mmol, 4 mol%), methyl acrylate (0.5 mL, 5.55 mmol) and Et₃N (1.5 mL, 11.1 mmol). The mixture was stirred at 80 °C for 21 h, cooled to r.t., and mixed with H₂O (40 mL). The organic compounds were extracted with EtOAc (50 mL), washed with brine (40 mL), dried (Na₂SO₄), and concentrated. The residue was purified by flash chromatography to provide 466 mg (47%) of recovered 1c followed by 450 mg (49%) of 2c (4,5-Ome).

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Preparation PP5

Heck Coupling of 1d and Methyl Acrylate to Form 2d (R = 5-NO₂): The procedure is same as that of 2c, yielding 82% of 2d after purification.

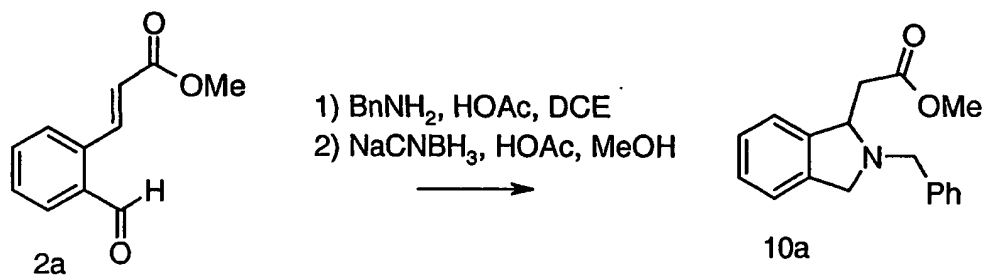
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Preparation PP6

Reductive Amination

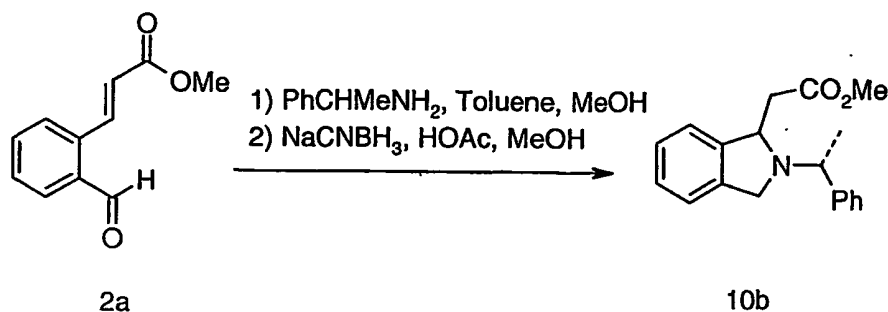


Reductive amination of (2a) with benzyl amine to form isoindoline (10a).

- 10 To a solution of 2a (11.27 g, 59.2 mmol) in ClCH₂CH₂Cl (60 mL) was added BnNH₂ (6.47 mL, 59.2 mmol), followed by HOAc (5.1 mL, 89 mmol). The mixture was stirred at r.t. for 1 h. NaCNBH₃ (5.58 g, 88.8 mmol) and MeOH (30 mL) were then added to the above solution. The resulting mixture was stirred at r.t. for another 2 h and quenched with sat. NaHCO₃ solution (150 mL). The mixture was extracted with EtOAc (2 × 100
- 15 mL) and the combined organic layers were washed with brine (150 mL), dried (Na₂SO₄), and concentrated to provide 15.3 g of crude product of 10a which was carried out for the next hydrogenolysis reaction.

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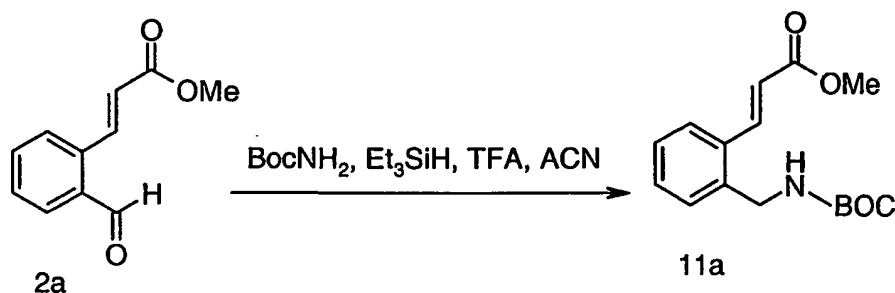
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Preparation PP7

One-pot process from 2-carboxaldehyde-methyl-cinnamate to target cyclized isoindoline product using NaBH_3CN . 2-carboxaldehyde-methyl-cinnamate 2a (3.254 g, 17.1 mmol) was dissolved in a 1:1 MeOH: PhCH_3 mixture (20 mL) at r.t. R-(+)-phenethylamine (2.073 g, 17.1 mmol) was added and the solution was heated under reflux for 2 hours. HPLC in process control indicated that the imine formation was completed. Then, AcOH (2.055 g, 34.2 mmol) and NaBH_3CN (2.15 g, 34.2 mmol) were successively added at r.t., the reaction mixture being cooled with a water-bath. The reaction mixture was post-agitated overnight. Water (10 mL), MeOH (20 mL) and 37% HCl (2.8 mL) were successively added and the organic layer was extracted. The aqueous layer was washed with PhCH_3 (10 mL). Then, the aqueous layer was made basic with 5N NaOH (20 mL) and MeOH was concentrated to partly remove MeOH. Extraction with EtOAc (2X 25 mL) was performed. The combined organic layers were dried over MgSO_4 , filtered and rinsed with EtOAc (10 mL). The filtrates were concentrated under reduced pressure and the residual oil was dried under vacuum overnight at r.t. to afford the target cyclized isoindoline product 10b with 92% yield (4.642 g, 15.7 mmol). HPLC % area indicated that the 2 diastereomers were produced in a 55:45 ratio. ^1H NMR confirmed this result by integration of the methyl group of the phenethyl substituent. Note: The Heck or Heck-type coupling was performed in toluene with a slight excess of methylacrylate which was removed by distillation before the MeOH and the R-(+)-phenethylamine addition.

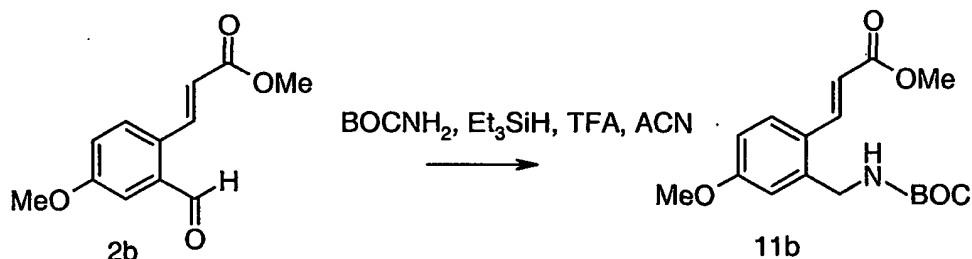
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Preparation PP8

Reductive amination of (2a) with *t*-butyl carbamate to form (11a): To a solution of aldehyde 2a (238 mg, 1.25 mmol) in CH_3CN (8 mL) was added *t*-butyl carbamate (439 mg, 3.75 mmol), followed by triethylsilane (0.6 mL, 3.75 mmol) and TFA (0.19 mL, 2.5 mmol). The mixture was stirred at r.t. overnight, quenched with sat. NaHCO_3 solution (20 mL) and extracted with EtOAc (2×30 mL). The combined organic layers were washed with brine (30 mL), dried (Na_2SO_4) and concentrated. The residue was purified by flash chromatography (hexane/EtOAc 3:1) to provide 317 mg (87%) of 11a.

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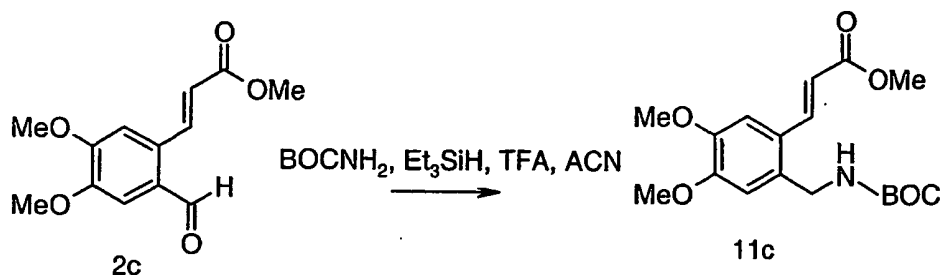
Preparation PP9

Reductive amination of 2b with *t*-butyl carbamate to form 11b: A mixture of aldehyde 2b (600 mg, 2.72 mmol) Et_3SiH (955 mg, 3 eq, 1.31 mL), TFA (620 mg, 2 eq, 420 μL), *t*-butyl carbamate (980 mg, 3 eq) in acetonitrile (15 mL) was stirred at room temperature over 2 days. Removed the solvent on a Rotary evaporator and purified the crude residue on a flash column (100 g SiO_2 , 7:1 \rightarrow 6:1 Hex / EtOAc). Collected 307 mg good desired product 11b (35 %); 195 mg product contaminated with aldehyde SM (22 %).

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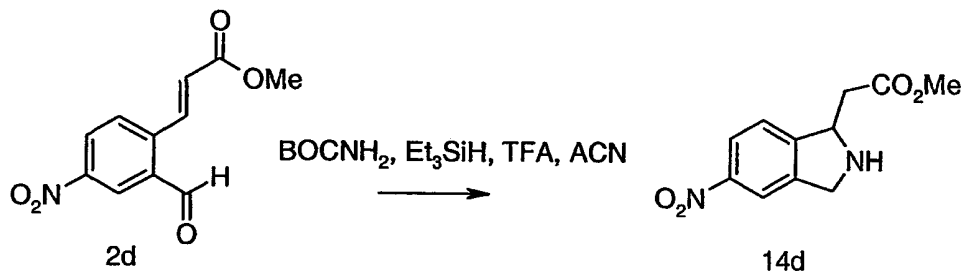
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Preparation PP10

Reductive amination of (2c) with *t*-butyl carbamate to form (11c): To a solution of aldehyde 2c (411 mg, 1.64 mmol) in CH₃CN (10 mL) was added *t*-butyl carbamate (580 mg, 4.93 mmol), followed by triethylsilane (0.8 mL, 4.93 mmol) and TFA (0.25 mL, 3.28 mmol). The mixture was stirred at r.t. overnight, quenched with sat. NaHCO₃ solution (30 mL) and extracted with EtOAc (2 × 30 mL). The combined organic layers were washed with brine (30 mL), dried (Na₂SO₄) and concentrated. The residue was purified by flash chromatography (hexane/EtOAc 3:1, hexane/EtOAc 1:1) to provide 535 mg (93%) of 11c.

15

Preparation PP11

To a solution of 2d (1.02 g, 4.34 mmol) in CH₂Cl₂/CH₃CN (1:1 24 mL) was added BocNH₂ (1.5 g, 13.02 mmol), Et₃SiH (2.1 mL, 13.02 mmol), and TFA (0.67 mL, 8.67 mmol). The mixture was stirred at r.t. for 7 h. A precipitate was formed during the reaction. The reaction mixture was quenched with sat. NaHCO₃ solution (30 mL), and diluted with CH₂Cl₂ (40 mL). The organic layer was washed with brine (30 mL), dried (Na₂SO₄), and concentrated. The residue was purified by flash chromatography (hexane/EtOAc 3:1, then CH₂Cl₂/EtOAc 10:1) to provide 2.08 g yellow solid which still

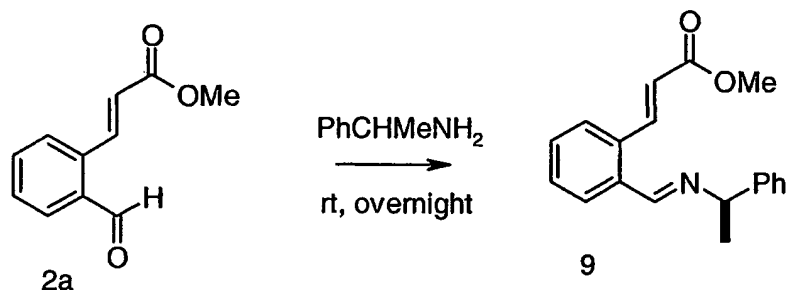
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- 5 containing BocNH_2 . The product is not the desired Boc-carbamate 14c. LC-MS result showed that the product is the Schiff base intermediate.

To the above product (420 mg) in CH_2Cl_2 (10 mL) was added Et_3SiH (1 mL) and TFA (0.4 mL). The mixture was stirred at r.t. for 1 h and small amount of sample was taken for NMR. NMR analysis demonstrated that the starting material was consumed and
10 the product was 14c. TFA (0.7 mL) was then added to the above mixture and the resultant solution was stirred at r.t. for another 5 h and concentrated. The residue was dissolved in EtOAc (20 mL) and washed with H_2O (10 mL). The aqueous layer was basified with sat. NaHCO_3 (30 mL) and the organic compounds were extracted with CH_2Cl_2 (2 X 25 mL). The combined organic layers were washed with brine (20 mL),
15 dried (Na_2SO_4) and concentrated to provide 218 mg of the cyclized compound 14c.

Preparation PP12



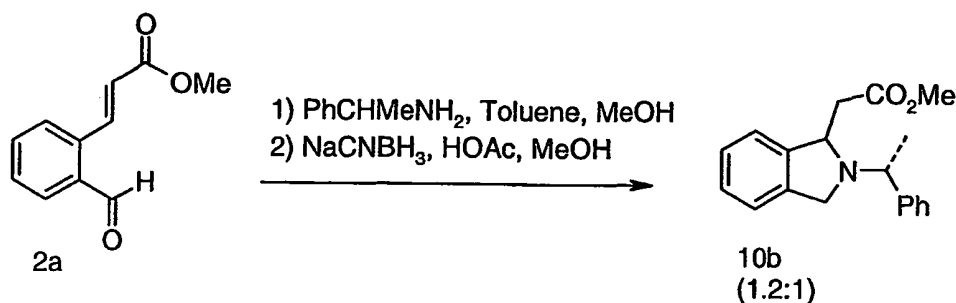
- 20 Condensation of 2a with alpha-methylbenzylamine to Form Imine 9. 2-carboxyaldehyde-methyl-cinnamate 2a (0.897 g, 4.72 mmol) was dissolved in MeOH (10 mL) at r.t. R-(+)-phenethylamine (0.577 g, 4.76 mmol) was added and the solution was heated under reflux for 2 hours. HPLC in process control indicated that the imine formation was completed. The solvent was stripped on a rotary evaporator and the
25 resulting oil was dried at r.t. under vacuum overnight. The Schiff base 9 was obtained almost quantitatively (1.412 g, 4.81 mmol).

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Preparation PP13

Michael Addition:



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The compound of α -methyl benzylamine was applied as the auxiliary. As shown above, the one-pot reaction of aldehyde **2a** and α -methyl benzylamine gave 90% of **10b** with a ratio of 1.2:1.

Step-wise reduction, amination, and cyclization:

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Condensation of aldehyde **2a** with α -methylbenzylamine in acetonitrile, methanol, methanol/toluene(1:1) or toluene afforded imine **9** in excellent yield. Reduction of the imine was initially carried out at r.t. with $\text{NaCNBH}_3/\text{HOAc}$. As a result, a poor ee ratio (1.2:1) was obtained, similarly to the previous described one-pot procedure. But when the reaction was carried out with NaBH_4/TFA at r.t., the ratio was elevated to 2:1. By

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lowering the reaction temperature to -78°C , the ratio was increased to 5 to 6:1.

Preparation PP14

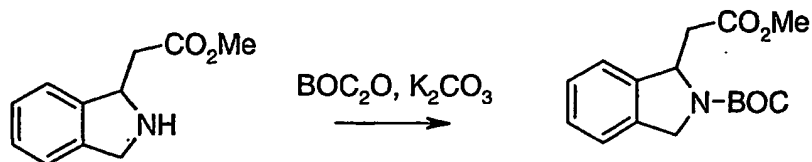
Cyclization of *t*-Butyl carbamate (**11a**): The *N*-Boc isoindoline methyl ester **12** was originally synthesized from **11a** via deprotection of Boc with TFA, followed by basic workup, and protection with a Boc group. This procedure has been greatly improved by a

25

one-step procedure.

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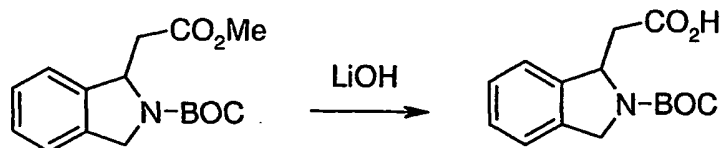
Preparation PP15

- In a 3 L 3-neck round bottom flask equipped with a nitrogen inlet,
- 10 thermocouple and mechanical stirrer, a solution of 160 g (1.15 moles) of K_2CO_3 in 180 mL of water was stirred at r.t. Solid BOC anhydride 120 g (0.55 moles) was added in one portion forming a semi-solution. To the reaction mixture, a solution of the crude amino ester starting material, 87 g (0.46 moles) in 120 mL of THF was added slowly at such a rate to keep the internal temperature below 35°C . A mild effervescence was observed.
- 15 The reaction mixture was stirred for 18 hours at r.t. Analysis of a reaction aliquot via NMR (DMSO-d_6) indicates the desired product. The reaction was diluted with brine and the product extracted with EtOAc. The organic layer was dried over Na_2SO_4 , filtered, and concentrated to yield a dark oil, 150.1 g, >100% yield. The crude material was taken on to the next step.

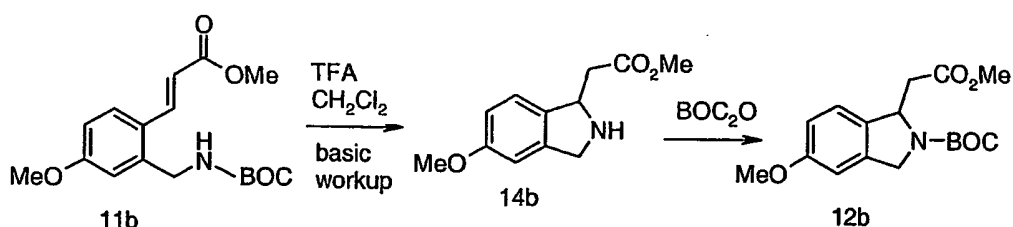
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Preparation PP16

In a 3-L 3-neck round bottom flask equipped with a mechanical stirrer, thermocouple, and reflux condenser, a solution of 150 g (approx. 0.46 moles) of crude N-BOC ester starting material in 750 mL of methanol was stirred at r.t. To the solution, 750 mL of water was added and the cloudy mixture was stirred vigorously. Solid LiOH 25 g (1.03 moles) was added in small portions at such a rate to maintain the internal temperature below 45 °C. Upon completion of addition, the reaction was stirred overnight at r.t. becoming a dark green color. After 18 hours the reaction was concentrated to yield a thick semisolid. The crude product was dissolved in EtOAc and washed with 1 N HCl quickly, followed by two brine washes. The organic layer was dried with Na₂SO₄, filtered and concentrated to yield 81 g of a dark green solid. The aqueous layers were combined and back extracted with methylene chloride, dried over Na₂SO₄, filtered, and concentrated to yield 6 g of a dark green solid. Both solids were combined to yield 87 g of desired product confirmed via NMR (DMSO₆).

Preparation PP17

Synthesis of 14b: Dissolved the N-boc compound 11b (200 mg, 0.62 mmol) in CH₂Cl₂ (1.0 mL). Cooled the clear light yellow solution to 0°C. Added slowly TFA (~710 mg, 10 eq, ~500 microliter) via a syringe. Removed the cooling bath and stirred the clear light brown solution at r.t. overnight. TLC (3:1 Hex / EtOAc, UV) confirmed a complete reaction. Removed the TFA on a rotavapor. Added EtOAc and concentrated again (twice). The crude residue was partitioned between EtOAc (10-15

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5 mL) and a sat. NaHCO_3 (10-15 mL). The aqueous was extracted with EtOAc (2x10 mL). The combined organic was dried over MgSO_4 , filtered, and concentrated to yield a light brown wet solid (212 mg, 138 %). NMR (CD_3OD) confirmed the desired isoindoline 14b. This crude isoindoline was used in the next protection step without purification.

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Preparation PP18

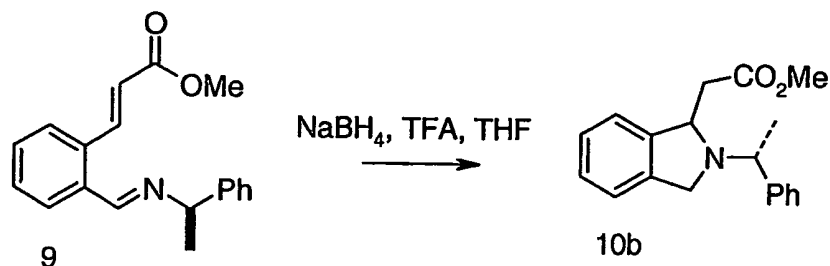
Synthesis of 12b: To a mixture of the isoindoline 14b (190 mg, 0.859 mmol), K_2CO_3 (189 mg, 1.5 eq) in a solvent 1:1 THF / H_2O (1.0 mL) at r.t. was added BOC₂O (210 mg, 1.1 eq). The reaction mixture was stirred at r.t. overnight. TLC (3:1 Hex / EtOAc, UV) confirmed a complete reaction. Diluted the mixture with EtOAc (15 mL), and washed with H_2O (1x20 mL). The aqueous was extracted with EtOAc (1x20 mL). The combined organic was washed with brine (1x20 mL), dried over MgSO_4 , filtered, concentrated to yield a clear brown oil (340 mg, 123 %). This crude oil was purified on a prep TLC plate (2x1,000 micron, solvent 2:1.5:0.5 CHCl_3 / Hex / EtOAc) to yield 12b a clear yellow oil (190 mg, 69 %). ^1H and ^{13}C NMR (CDCl_3) were obtained.

Procedure PP19

Synthesis of 12d (5- NO_2) by Boc-protection. The compound was prepared by following the same procedure as described for 12b.

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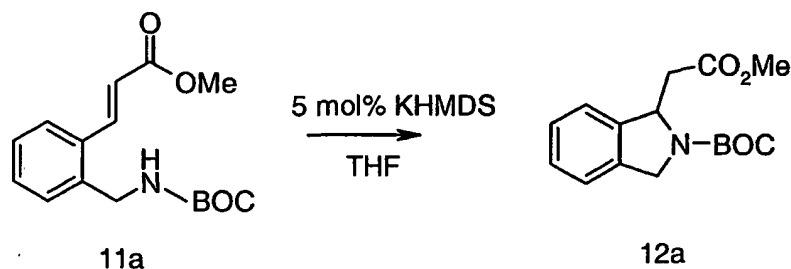
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Preparation PP20

The imine 9 (1.412 g, 4.81 mmol) was dissolved in anhydrous THF (10 mL) at r.t. and TFA (5 mL) was added. The black solution was then cooled to -78°C (dry ice bath) and NaBH₄ (0.893 g, 23.6 mmol, 5 eq.) was added in 2 portions over 5 minutes. Then, the reaction mixture was post-agitated at -78°C for 3 hours and allowed to gently warm at r.t. overnight. Water (20 mL), cyclohexane (10 mL) and EtOH (20 mL) were successively added and the organic layer was extracted and discarded. The aqueous layer was made basic with 5N NaOH (20 mL) and extracted two times with a 2:1 EtOAc/PhCH₃ mixture (30 mL). The combined organic layers were dried over MgSO₄, filtered and rinsed with EtOAc (10 mL). The filtrates were concentrated under reduced pressure and the residual oil was dried under vacuum overnight at r.t. to afford the target cyclized isoindoline product 10b (1.273 g, 4.31 mmol) with 91.4% yield. HPLC % area indicated that the 2 diastereomers were produced in a 84:16 ratio (de 68%). ¹H NMR confirmed this result by integration of the methyl group of the phenethyl substituent.

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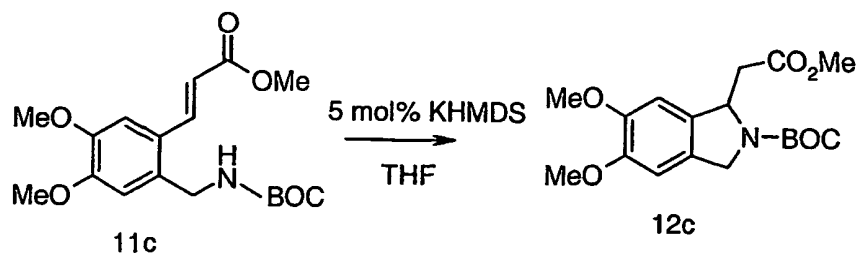
Preparation PP20

- N-Boc methyl ester 11a (36.3 g, 0.125 mol) was dissolved in THF (250 mL), and the solution was cooled to about 0°C. A solution of potassium bis(trimethylsilyl) amide (1.24 g, 0.05 mol. eq.) was added slowly via a syringe under nitrogen atmosphere. The temperature was raised about 8 degrees during the addition. The cooling bath was removed and the solution was stirred at r.t. for 30-45 min. The clear brown solution was poured into a separation funnel containing about 100 mL of a saturated NH₄Cl. The layers were separated. The aqueous layer was extracted with EtOAc (2x50 mL). The combined organic was washed with brine (1x100 mL), dried over Na₂SO₄, filtered, concentrated on a Rotary evaporator to a clear yellow oil (37.3 g). This crude oil was purified on a flash column (600 g SiO₂), with a gradient solvent 6:1 Hex/EtOAc (2.1 L), 5:1 Hex/EtOAc (1.2 L), 4:1 Hex/EtOAc (1.5 L) to yield 12a as a clean yellow oil (34.5 g, 95 %).

-200-

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Preparation PP21

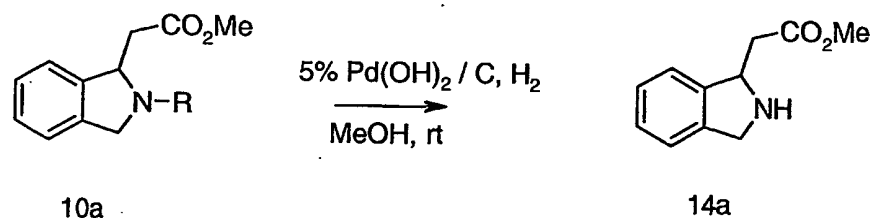


To a solution of 11c (535 mg, 1.52 mmol) in THF (10 mL) was added KHMDS (0.5 M in toluene, 0.1 mL, 0.05 mmol, 2 mol%). The mixture was stirred at r.t. for 20 min, quenched with sat. NH₄Cl solution (20 mL), and diluted with EtOAc (20 mL).

10 The organic layer was separated, washed with brine (20 mL), dried (Na_2SO_4) and concentrated. The residue was filtered through a plug of silica gel ($\text{EtOAc}/\text{CH}_2\text{Cl}_2$ 1:10) to give 530 mg (99%) of 12c as an off white solid.

Preparation PP22

15 Deprotections:

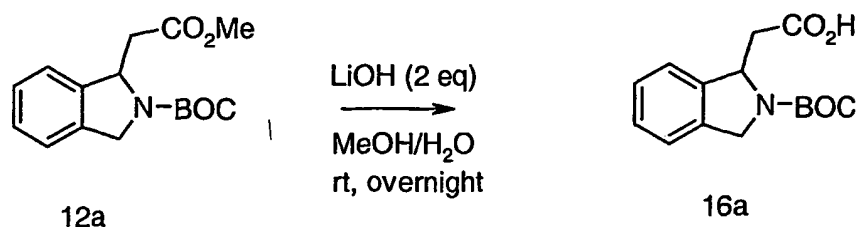


Hydrogenolysis of 10a (R = Bn) to Form (14a): To a solution of crude 10a (15.3 g, 54.4 mmol) in MeOH (100 mL) was added Pd(OH)₂/C (Pearlman's catalyst, 1.02 g, 6 mol%) in a par-shaker bottle. The suspension was shaken under 30 psi H₂ pressure overnight in the par-shaker, and filtered through a plug of celite. The filtrate was concentrated to provide 10.1 g of crude 14a as brown oil. (The procedure is same for the methyl benzylamine isoindoline substrate 10b)

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Preparation PP23

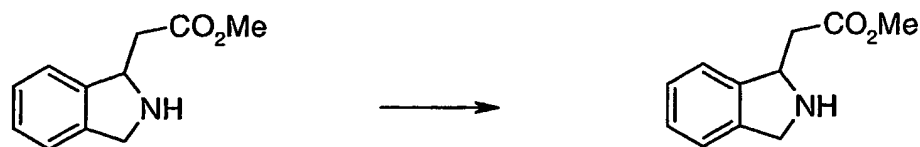
In a typical reaction a mixture of the isoindoline ester 12a (92 mg, 0.316 mmol) in 1:1 MeOH/H₂O (2 ml) was treated with LiOH (15 mg, 2 eq) at r.t. overnight. Diluted the mixture with CH₂Cl₂ (5 ml) and water (5 ml). Adjusted the pH of the reaction mixture to 1-3 with a 10% NaHSO₄ solution. Separated the layers. The aqueous was extracted with CH₂Cl₂ (1 x 10 ml). The combined organic was dried over Na₂SO₄, filtered, concentrated to yield 16a as a pale yellow foam (76 mg, 87 %). NMR (CDCl₃) showed a clean desired acid product.

It is noted that the reaction time must be more than 6 hours. The crude foam can be purified by slurry in warm hexane and then filter to yield a tan solid. Hydrolysis using KOH (2-5 eq) in 1:1 MeOH/H₂O overnight would give the same result.

20

Preparation PP24

Resolution:



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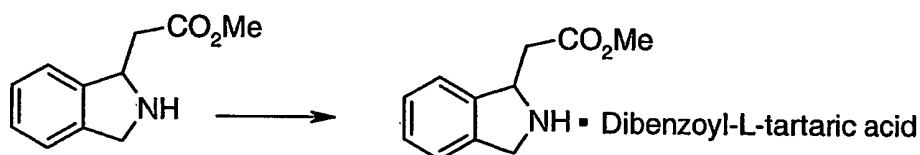
Purification of Partially Resolved Isoindoline-carboxylic acid methyl ester:

A solution of the crude material (97.62g) isoindolinecarboxylic acid methyl ester in CH₂Cl₂ (350 mL) was extracted with 1M HCl (400 mL, 200 mL). The combined aqueous portions were washed with CH₂Cl₂ (4 X 250 mL) and then made basic with K₂CO₃ solution (85 g in 150 mL of water). The mixture was extracted with CH₂Cl₂ (6 X 100

-202-

- 5 mL) and the combined organic extracts were dried (Na_2SO_4) and concentrated to give partially resolved Isoindolinecarboxylic acid methyl ester as an oil (33.2 g). 60% ee by chiral CE.

10 Preparation PP25

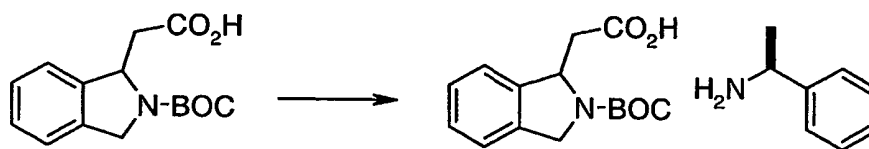


Resolution of Partially Resolved Isoindoline-carboxylic acid methyl ester:

- A solution of partially resolved isoindoline-carboxylic acid methyl ester (33.24 g, 0.174 mol) in EtOH (130 mL) was treated slowly with a solution of dibenzoyl-L-tartaric acid (56.06 g, 0.156 mol) in EtOH (200 mL). The solution was seeded with seeded with product and stirred at r.t. for 4 hours. Pure product was collected by filtration, washed with EtOH (30 mL) and dried to off-white crystals (60.49 g). 96.5% ee by chiral CE.

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Preparation PP26



Resolution of N-BOC Isoindolinecarboxylic acid: A solution/slurry of

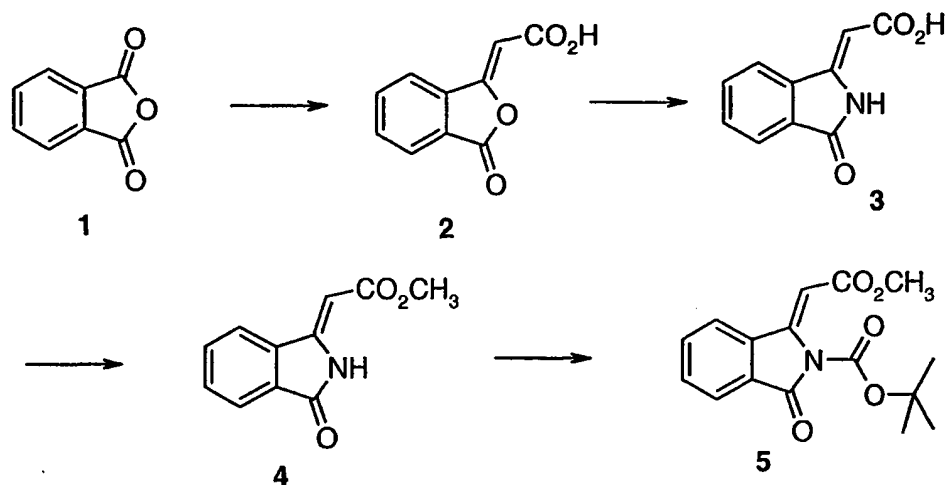
- 25 racemic N-BOC Isoindolinecarboxylic acid (114.5 g, 0.413 mol) in EtOAc (1000 mL) was treated slowly with triethylamine (28.8 mL, 0.206 mol), followed by (S)-(-)-alpha-methylbenzylamine. The solution was seeded with product and stirred at r.t. overnight. The product was collected by filtration, washed with EtOAc (200 mL) and dried to a white powder (62.98 g). 97.6% ee by chiral CE.

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5 Asymmetric Hydrogenation Routes

Part I: Synthesis of the Z-isomer (precursor of asymmetric hydrogenation)

Scheme P1Preparation PP27

Z-isomer 5 was synthesized as outlined in Scheme P1. Compound 5 was shown to be a single isomer by HPLC and H-1 nmr. The double bond stereochemistry was derived from comparative NOE data using the purported E-isomer (Scheme P1). The best chiral induction was achieved using compound 8/Ferrotane/MeOH-THF. With regard to the conversion of 9 to 10, which would constitute a formal asymmetric synthesis of isoindolene 10, this has been achieved using Super hydride-BF₃.OEt₂. However, the product was a mixture of 10 and the corresponding de-BOC (deprotected) compound.

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Preparation PP28

Compound 2 (scheme P1)

- Phthalic anhydride (751.5g, 5.014 mole), potassium acetate (498g, 5.014 mole) and acetic anhydride (1L) were stirred together under nitrogen. The mixture was slowly warmed to 145-150°C and stirred for 10 minutes, then at 140°C for 20 minutes.
- 10 The mixture was allowed to slowly cool to 80°C over 1 hour. Three volumes of water were added causing precipitation of a solid. After filtration, the filtered solid was washed with warm water and pulled as dry as possible for 30 minutes. The solid was then washed with ethanol and acetone respectively. If required further purification could be achieved by slurrying the solid in acetone, at room temperature, for 15 minutes, then filtration.
- 15 Drying in vacuo at 50°C for 20 hours gave compound 2 as an off-white solid, 470g (48%) with an NMR purity of approx. 90%.

Preparation PP29

Compound 3 (Scheme P1)

- 20 Compound 2 (470g, 2.47 mole) was added to stirred aqueous ammonia (470 ml conc. NH₃ in 4.7L water). The resultant mixture was stirred at room temperature for 1 hour then filtered. The filtered solid was washed with water. The combined aqueous filtrate and washings were carefully acidified with 6M aq. HCl (2.35L). The precipitate was removed by filtration and dried in vacuo at 50°C to give compound 3 as a
- 25 yellow solid, 259g (52%).

Preparation PP30

Compound 4 (Scheme P1)

- Compound 3 (511g, 2.7 mole) was slurried in toluene (10 vol). Thionyl chloride (385g, 3.24 mole) was added over 10 minutes to the stirred mixture, which was
- 30 then heated to reflux for 1.5 hours. H-1 NMR analysis indicated approx. 80% conversion to acid chloride). DMF (3.7ml) was added and the mixture refluxed an additional 3 hours. The resultant mixture was allowed to cool to 35°C and methanol (1.27L) added at such a rate that the reaction temperature was maintained at 30-35°C. The reaction

-205-

- 5 mixture was kept at this temperature a further 15 minutes then concentrated in vacuo to give compound 4 as a brown solid, 536g (quantitative).

Preparation PP31

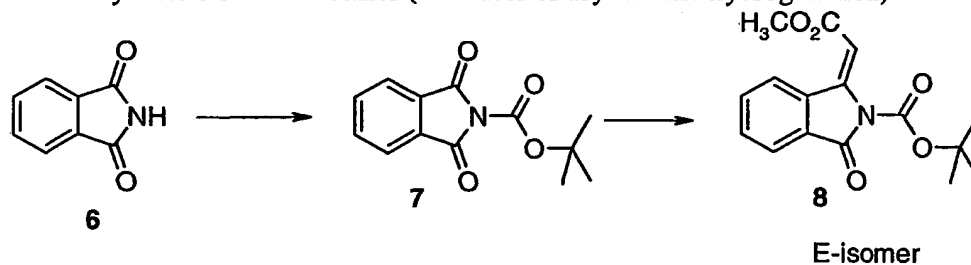
Compound 5 (Scheme P1)

- 10 Compound 4 (750g, 3.65 mole) was dissolved in acetonitrile (15L). The stirred mixture was cooled to 0-5°C and DMAP (624g, 5.11 mole) added in one portion. After 10 minutes BOC anhydride (1115g, 5.11 mole) was added in one portion: there was a slight exotherm accompanied by gas evolution. The mixture was stirred at room temperature for 5 hours, and then concentrated in vacuo. The residue was dissolved in
- 15 EtOAc and washed with 10% aqueous citric acid, satd. aq. Na₂CO₃ and water respectively. After drying, concentration of the organics gave a thick syrup. This material was run through a plug of silica gel (1.5kg) eluting with 1:1 EtOAc-hexane. Compound 5 was isolated as a dark solid, 619g (55%). Careful chromatography on silica gel eluting with 20% EtOAc-hexane gave 5 as a fluffy white solid.

20

Scheme P2

Part II: Synthesis of the E-isomer (Precursor of asymmetric hydrogenation)



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Preparation PP32

The E-isomer of Compound 8 (Scheme P2) was prepared as shown in Scheme P2.

10

Preparation PP33

Compound 7 (Scheme P2)

The compound 7 was prepared according to the procedure of Einhorn et al, *Synth. Commun.* 2001, 31(5), 741-748.

15

Preparation PP34

Compound 8 (Scheme P2)

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Compound 7 (15.00g, 60.7 mmole) and methyl(triphenyl phosphoranylidene) acetate (41.40g, 121.3 mmole) were slurred in toluene (150ml). The mixture was stirred at reflux and monitored for reaction of 7 by GC. After 1.5 hours the reaction appears complete by GC. After cooling to room temperature, the mixture was filtered. The solid on the filter was washed with toluene until colorless. The combined filtrate / washings were concentrated in vacuo to leave a tan solid. This material was coated on silica gel and chromatographed on silica gel (1kg) eluting with 10% EtOAc-hexane. Compound 8 was isolated as a white or pale yellow powder, 5.52g (30%).

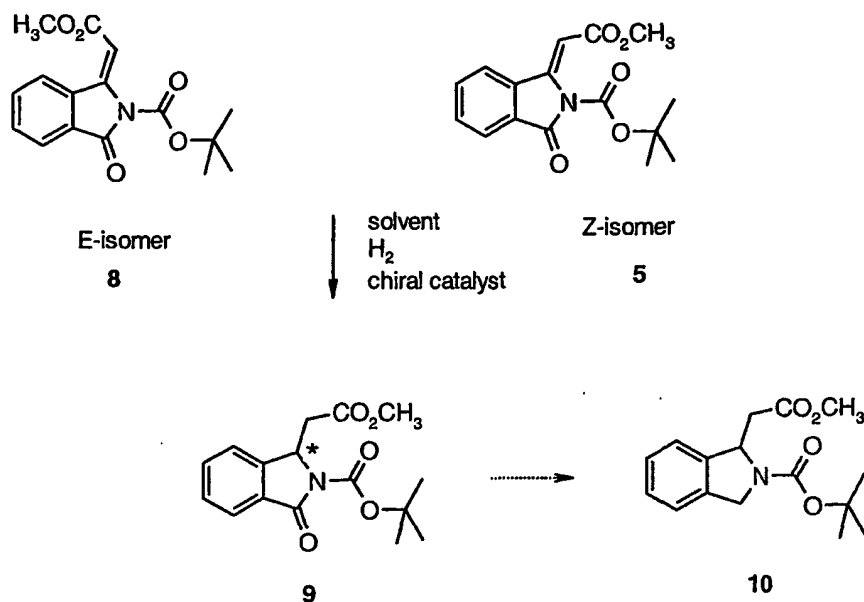
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Scheme P3

Asymmetric hydrogenation:



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Preparation PP35

Screening of chiral hydrogenation conditions indicated that the best chiral induction was achieved using compound 8/Ferrotane/MeOH-THF. With regard to the conversion of 9 to 10, which would constitute a formal asymmetric synthesis of isoindolene 10, this has been achieved using Super hydride- $\text{BF}_3 \cdot \text{OEt}_2$. However, the product was a mixture of 10 and the corresponding de-BOC (deprotected) compound.

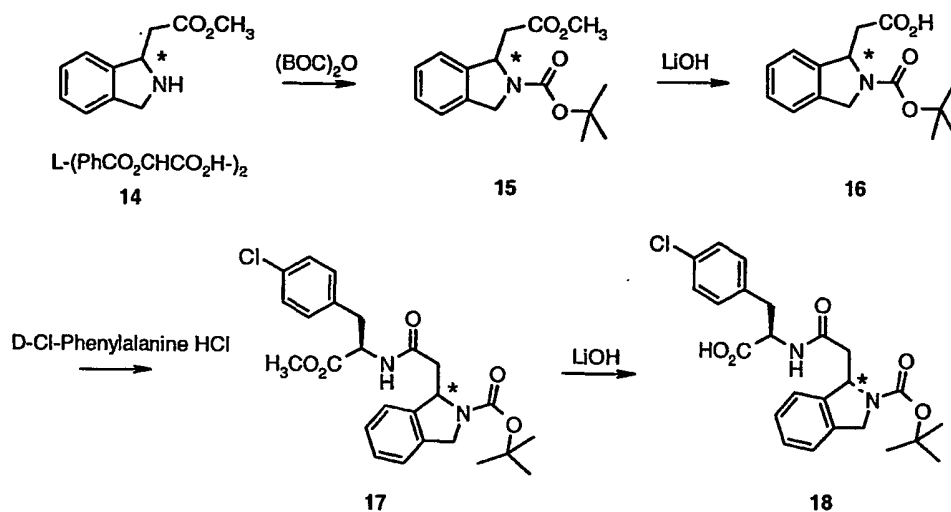
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Scheme P4

Coupling of chiral isoindoline with d-4-chloro-Phenyl-alanine using tartrate salt:



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Preparation PP36

Compound 15 (Scheme P4)

Tartrate salt 14 (58.00g, 100.27 mmole) was slurred in water (580ml). Solid NaHCO₃ (25.27g, 300.8 mmole) was carefully added. BOC anhydride (22.98g, 105.28 mmole) was added in one portion and the progress of the reaction monitored by reverse phase HPLC. After 1 hour additional BOC anhydride (2.18g, 10.00 mmole) was added. The reaction was complete (by HPLC) after 3 hours. The mixture was extracted with EtOAc (2 x 250ml). The combined organic extracts were washed with water (250ml) and dried (MgSO₄). Filtration and concentration in vacuo gave 15 as a clear light brown oil (31.33g) contaminated with a small amount of t-BuOH and BOC anhydride.

This material was used directly in the next reaction.

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Preparation PP37

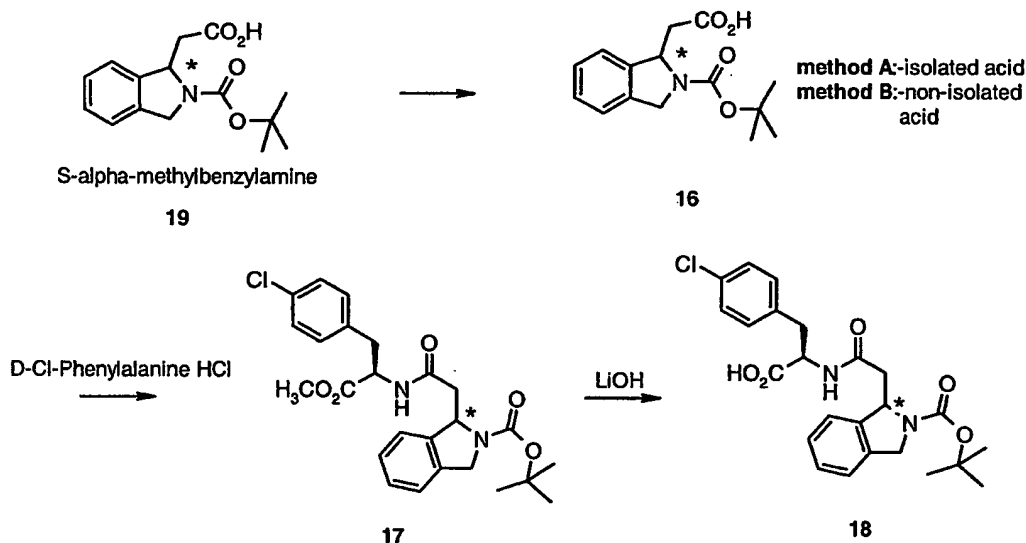
Compound 16 (Scheme P4)

- Ester 15 (29.21g, 100.26 mmole) was dissolved in 3:1 THF-water (100ml). LiOH (6.00g, 250.65 mmole) was added in 1 portion to the stirred solution. After 17 hours, the mixture was stripped to dryness and the residue dissolved in water (500ml).
 10 EtOAc (250ml) was added and solid NaHSO₄ added to the stirred mixture until the pH=3. The organic layer was separated and the aqueous layer extracted with EtOAc (250ml). The combined EtOAc layers were dried (MgSO₄). Filtration and concentration in vacuo gave acid 16 as a light tan solid, 27.10g (97%).

15

Scheme P5

Rrom alpha-methyl benzylamine salt:



20

The chemistry used is shown in Scheme P5. Two protocols were used: method A used isolated 16, method B used a solution of 16 derived from resolved salt 19.

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Preparation PP38

Compound 17 (Scheme P5, method A)

Acid 16 (24.18g, 87.2 mmole) and D-chloro-phenylalanine hydrochloride (21.81g, 87.2 mmole) were dissolved in CH₂Cl₂ (100ml) and DMF (25ml). The mixture was stirred at ambient temperature. HOBt (13.55g, 100.3 mmole) and Hunig's base (45.6ml, 33.81g, 261.6 mmole) were added. HATU (38.13g, 100.3mmole) was added in 10 1 portion (there was a rapid exotherm to 50°C). The mixture was stirred for 90 minutes then diluted with EtOAc (750ml). The resulting mixture was washed with water, 5% KHSO₄, brine and satd. NaHCO₃ respectively, then dried. Filtration and concentration in vacuo gave crude 17 as a brown foam. The product was purified by chromatography on 15 silica gel (1kg) eluting with 1:1 EtOAc-hexane. Ester 17 was isolated as a tan powder, 38.85g (94%).

Preparation PP39

Compound 17 (Scheme P5, method B)

20 Resolved salt 19 (96.27g, 232.5 mmole) was partitioned between water (500ml) and CH₂Cl₂ (250ml) Solid KHSO₄ was added portion wise until pH=2.5. Separate the organic layer and extract the aqueous layer with CH₂Cl₂ (150ml). The combined organic layers were dried (MgSO₄) then filtered. To this solution was added 4-chloro-D-phenylalanine (58.16g, 232.5 mmole), HOBt (34.57g, 255.8 mmole), Hunig's 25 base (93.2ml, 69.13g, 534.9 mmole) and finally HATU (97.26g, 255.8 mmole). The resultant mixture was stirred at room temperature for 18.5 hours, and then poured onto a plug of silica gel (1kg). This was washed with 1:1 EtOAc-hexane until no more product elutes. Ester 17 was isolated as a pink foam, 101.79g (93%): contains about 1% unreacted 16.

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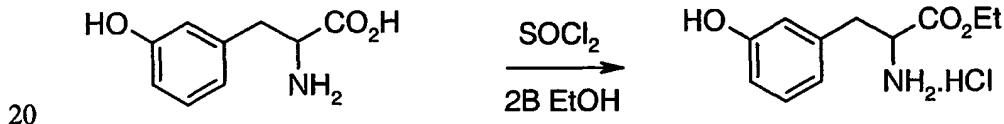
-211-

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Preparation PP40

Compound 18 (Scheme P5)

Ester 17 (38.64g, 81.7 mmole) was dissolved in 3:1 THF-water (200ml). LiOH (2.15g, 89.9 mmole) was added to the mixture, which was stirred at room temperature for 2 hours. The solvent was then removed in vacuo and the residual solid taken up in water (600ml). This was extracted with MTBE (250ml). The aqueous layer was separated and stirred with EtOAc (250ml), and solid KHSO₄ was added portion wise until pH=3. The layers were separated and the aqueous extracted with EtOAc (250ml). The combined organic layers were dried over MgSO₄. Filtration and concentration in vacuo gave acid 18 as a light pink foam, 38.41g (35.71g corrected for residual solvent, 15 95%).

Preparation PP41Step 1: Esterification

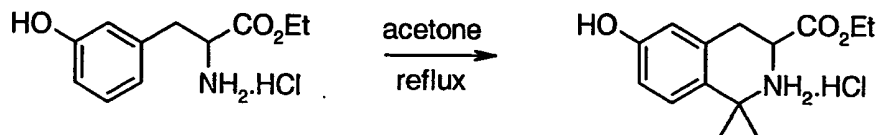
In a 22 L 4-neck round bottom flask equipped with a reflux condenser, thermocouple and nitrogen inlet, a slurry of 1000 g (5.4 moles) of m-tyrosine in 10 L of 2B-3 EtOH was cooled to 5 °C. To the slurry, 350 mL (12.4 moles) of thionyl chloride were added dropwise via an addition funnel at such a rate to maintain the reaction temperature below 20 °C. Upon completion of addition, the reaction was heated to reflux temperature and stirred for 18 hrs. The reaction was concentrated to one-third the volume and 8 L of MTBE were charged. The resulting thick slurry was stirred for 14 hrs in a rotary evaporator at r.t.. The resulting solid was isolated on a filter pad and dried at 40 °C for 48 hrs yielding 1288 g (95%). NMR (DMSO-d₆) indicated desired material.

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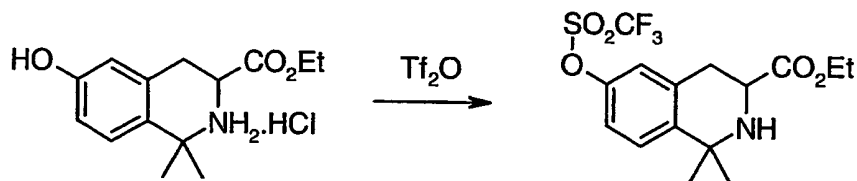
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Preparation PP42Step 2: Pictet-Spengler

In a 22 L 4 neck round bottom flask equipped with a mechanical stirrer,
 10 thermocouple, and reflux condenser placed on top of a Soxhlet extractor charged with 4
 °A sieves, a semi-solution of *m*-tyrosine ethyl ester hydrochloride 1288 g (5.26 moles) in
 13 L of acetone was heated to reflux temperature. The condensate was filtered through
 the sieves to remove water. The reaction was stirred vigorously at reflux for 48 hrs. An
 NMR sample in DMSO_d₆ indicated the absence of starting material. The reaction was
 15 cooled to r.t. and concentrated to yield an off-white solid, 1411 g (94%).

Preparation PP43Step 3: Triflation

20

In a 22 L 4 neck round bottom flask equipped with a reflux condenser,
 mechanical stirrer, nitrogen inlet, and a thermocouple, 1240 g (4.35 moles) of the starting
 material salt in 12.4 L of methylene chloride was cooled to 4 °C. To the mixture, 1452
 25 mL (10.4 moles) of triethylamine were added and stirred into solution. Triflic anhydride,
 1472 mL (5.22 moles) was added dropwise to the reaction at such a rate to maintain the
 internal temperature below 10 °C. The ice bath was removed and the reaction warmed to
 r.t. and stirred for 18 hrs. The reaction was concentrated to a oil then dissolved in 4 L of
 EtOAc and concentrated again to an oil in an effort to remove excess triflic anhydride
 30 The crude residue was dissolved in 4 L of EtOAc and washed with water and saturated
 sodium bicarbonate solution. The organic layer was isolated and dried with sodium

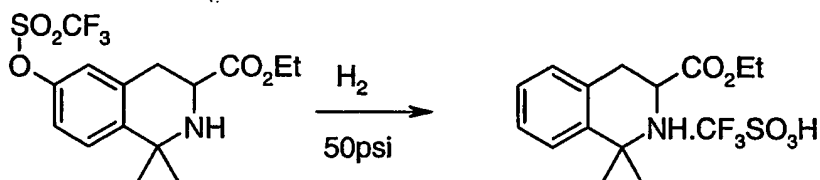
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- 5 sulfate, filtered and concentrated to yield 1720 g (>100%) of a crude dark oil which was used without further purification.

Preparation PP44

Step 4: Deoxygenation

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- A solution of 1720 g (4.35 moles) of crude starting material in 14 L of acetone was charged to a 10 gallon stainless steel autoclave. To the solution, a slurry of 5% Pd/C in 1.2 L of toluene was added. The reaction mixture was evacuated and purged with H_2 gas at 50 psi two times. The reaction was stirred overnight at 50 °C with H_2 at 50 psi. A sample aliquot indicated no reaction had occurred. The mixture was filtered and concentrated to a thick oil and resubjected to reaction conditions. After 18 hrs, NMR of a sample aliquot indicated absence of starting material. The reaction mixture was filtered and the filtrate concentrated to yield 1581 g of an off-white solid (95%).

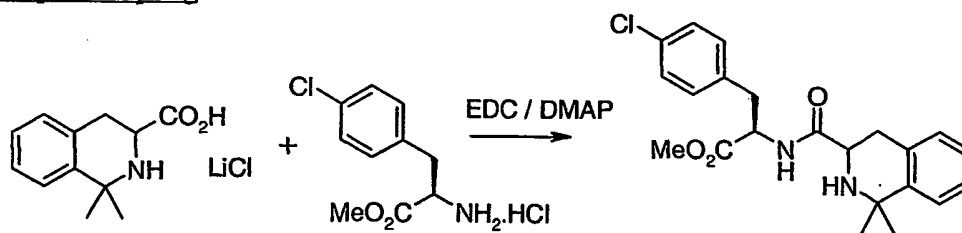
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Preparation PP45Step 5: Hydrolysis/Salt Formation

To a 2 L 3 neck round bottom flask equipped with a mechanical stirrer,
 10 thermocouple, and nitrogen inlet, a mixture of 700 g (1.83 moles) of the triflate salt
 starting material was charged. A solution of 427 g (1.83 moles) of the starting material
 free base in 13.3 L of THF was added followed by 700 mL of water. The semi-solution
 was stirred vigorously at r.t. To the reaction flask, 43.7 g (1.83 moles) of solid LiOH
 were added in small portions at such a rate to maintain the internal temperature below 35
 15 °C. The reaction was stirred for 18 hrs at r.t. and concentrated to yield a thick oil. THF
 (4 L) was added and the semi-solution was concentrated. This was repeated with toluene
 and the semi-solid was placed under house vacuum on the roto vap with stirring for 18 hrs
 to yield 650 g of a crude solid. The solid was reslurred in EtOAc, filtered, isolated and
 dried to yield 525 g (68%) of the lithium salt as an off-white solid.

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Preparation PP46Step 6: Coupling

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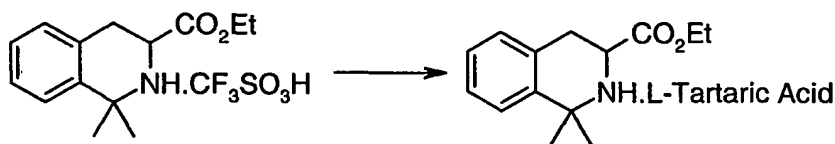
Solid *d*-chloro-phenylalanine 446 g (1.78 moles) was added to the semi-
 solution followed by 20 g (0.162 moles) of DMAP. The resulting mixture was stirred for
 15 minutes then solid EDCI (1-(3-dimethylaminopropyl)-3-ethylcarbodiimide
 hydrochloride) 390 g (2.03 moles) was added. The reaction mixture was heated to 80 °C
 and stirred for 18 hours. Thin layer chromatography (1:1 EtOAc:Hex) indicated very

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- 5 little starting material present. The reaction was cooled to r.t. and concentrated to yield a thick oil. The crude oil was dissolved in EtOAc and washed with water, and brine. The solution was dried with sodium sulfate, filtered and concentrated to yield a thick oil, 426 g. The crude oil was chromatographed in several lots using a Waters Prep 500 chromatography apparatus. The eluent consisted of a gradient system, 5%-80% EtOAc in
10 heptane at a flow rate of 240 ml/min over 38 minutes. The two diastereomers were separated and isolated to yield 119.04 g for the top spot and 111.3 g for the bottom spot. Conformation of both desired diastereomers was achieved via NMR (DMSO₆).

Preparation PP47

- 15 Resolution of tetrahydroisoquinolinecarboxylic acid ethyl ester to prepare l-tartaric acid salt:



- 20 Preparation of free-base: A racemic mixture of tetrahydroisoquinolinecarboxylic acid (7.43 g) in EtOAc (60 mL) was treated with saturated NaHCO₃ solution (60 mL) and saturated Na₂CO₃ solution (10 mL). The mixture was agitated and the layers were separated. The organic phase was dried (Na₂SO₄) and concentrated to give the corresponding free-base as an oil (4.85 g)
- 25 Resolution: A mixture of the above free base (467 mg, 2.0 mmol), and L-tartaric acid (300 mg, 2.0 mmol) in acetone (4 mL) was stirred at r.t. overnight. The title L-tartaric acid salt was collected by filtration, washed with acetone (about 2 mL) and dried to a white powder (367 mg). 100% ee by chiral CE.

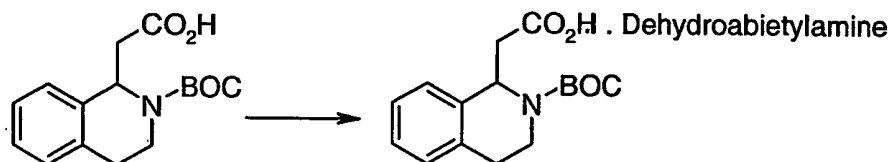
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Preparation PP48

Resolution of N-BOC tetrahydroisoquinolinecarboxylic acid



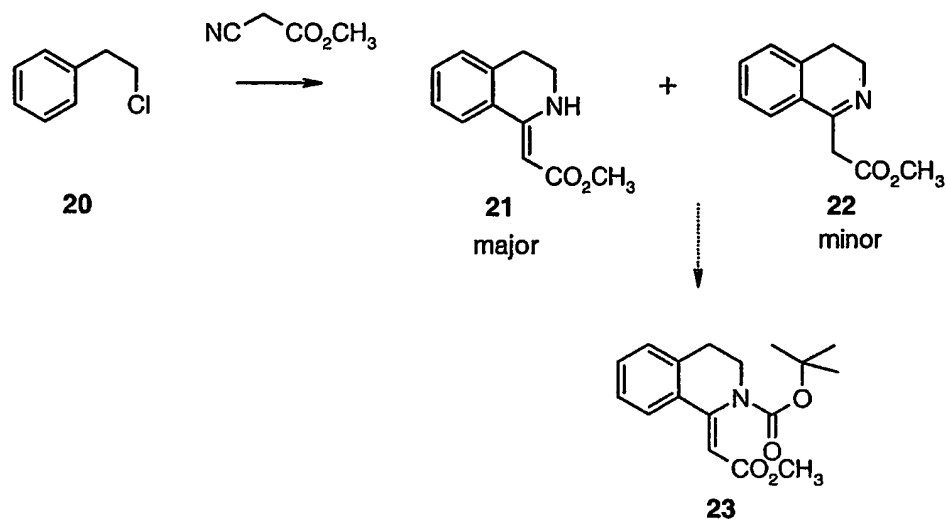
- 10 2-{2-[(tert-butyl)oxycarbonyl]-1,2,3,4-tetrahydro-isoquinolyl}acetic acid
 dehydroabietylamine salt: Racemic 2-{2-[(tert-butyl)oxycarbonyl]-1,2,3,4-
 tetrahydroisoquinolyl}acetic acid (30.15 g, 103.5 mmol) was dissolved in *i*-PA (300 mL).
 Dehydroabietylamine (22.11 g, 52.7 mmol of a 68 weight % mixture) was added to the
 solution, which was then agitated on a multi-arm shaker for 63 h. The resultant thick
 15 paste was filtered and rinsed with *i*-PA (50 mL, 25 mL). Dried in a 50 °C vacuum oven
 to obtain a white solid (27.73 g, 52 % ee by chiral CE analysis). The product was
 reslurred in *i*-PA (266 mL) and agitated on a multi-arm shaker for 23.5 h. Filtered the
 thick slurry and rinsed with cold *i*-PA (50 mL, 30 mL). Dried the cake in a 50 °C vacuum
 oven and obtained the product as a white solid (23.63 g, 40 % yield, 94 % ee by chiral CE
 20 analysis).

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Scheme P6

Asymmetric Hydrogenation:



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Preparation PP49

Enamine 21 (Scheme P6) was prepared as a substrate for asymmetric hydrogenation screening studies. It is formed as an approx. 10:1 mixture with imine 22. The enamine (21) may be NH-protected i.e., by a Boc protecting group. The resulting compound 23 may be subjected to asymmetric hydrogenation to afford the acetic acid or methylacetate substituted isoquinoline, which may be processed into a compound of formula I as demonstrated previously.

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Preparation PP50

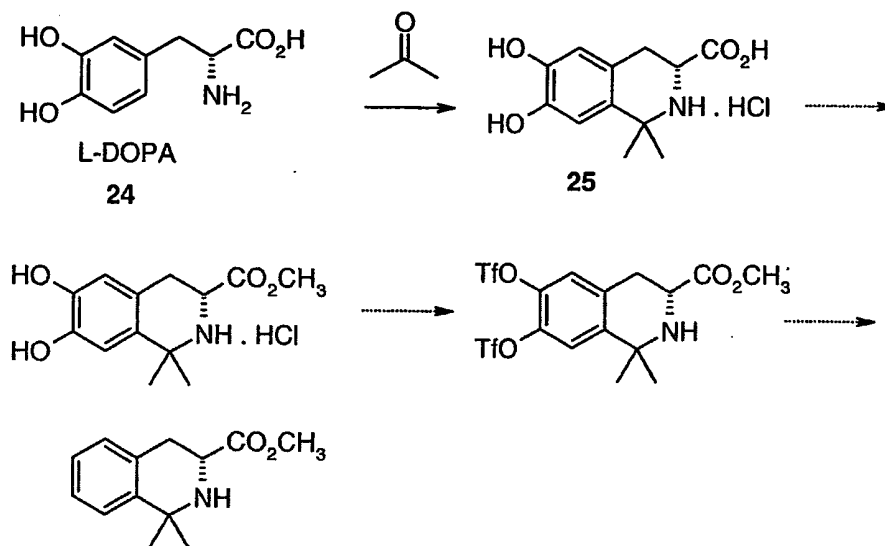
Compound 21 (Scheme P6)

Prepared as published W Sobotka et al, *J. Org. Chem.*, 1965, 30, 3667.

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Scheme P7

Synthesis of Gem-dimethyl TIC:

Preparation PP51

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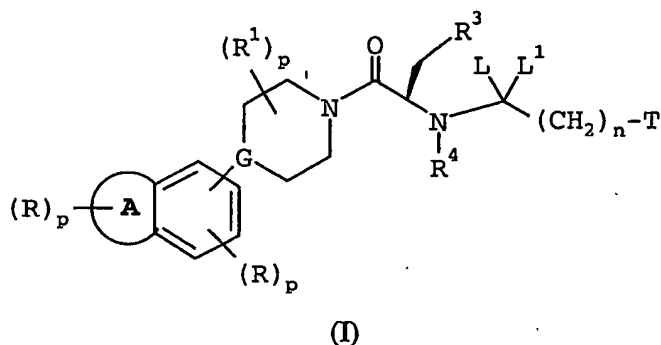
The chiral synthesis of gem-dimethyl TIC using L-Dopa as the starting material instead of tyrosine was successfully demonstrated up to the Pictet-Spengler reaction with L-DOPA and acetone. The product is a mixture of starting material 24 and product 25 (major component). The product was isolated by using common isolation procedures. An alternative isolation method is to react the mixture (24 and 25) with BOC anhydride wherein the less hindered N-H in 24 leads to preferential BOC protection of 24, allowing for ready separation of 25. Chemistry for the rest of the sequence i.e., deoxygenation reaction, has been demonstrated herein.

25

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5 WHAT IS CLAIMED IS:

1. A compound of formula I:



10

or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

G is CR¹ or N;

- 15 L and L
- ¹
- are independently hydrogen or together oxo;

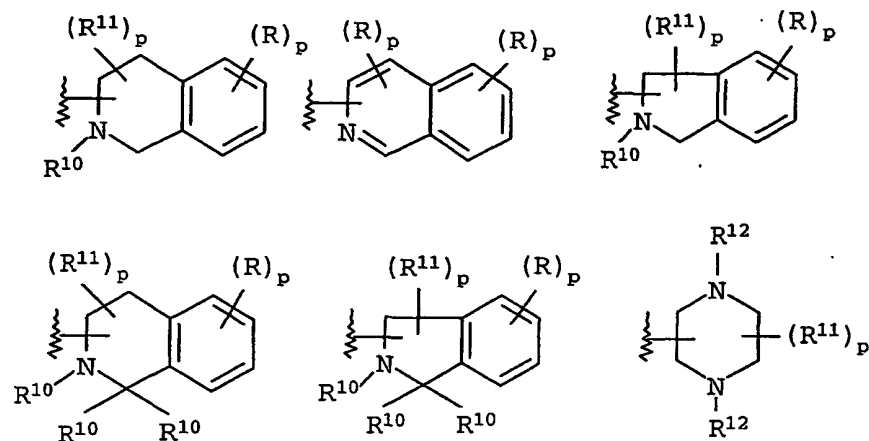


A is a 5- or 6-membered carbocyclyl wherein the carbocyclyl being saturated, partially saturated or aromatic ring, and the carbocyclyl being optionally substituted with one to three substituents independently selected from R;

20

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5 T is:



R is independently:

- hydrogen,
- hydroxy,
- 10 (D)cyano,
- halo,
- C₁-C₈ alkyl,
- C₁-C₈ alkoxy,
- C₃-C₇ cycloalkyl,
- 15 C₁-C₄ haloalkyl,
- (D)heterocyclyl
- (D)C(O)R⁸,
- (D)C(O)(CH₂)_nN(R⁸)₂,
- C₁-C₈ alkyl-N(R⁸)₂,
- 20 (D)OR⁸,
- (D)OCOR⁸,
- (D)OC(O)N(R⁸)₂,
- (D)N(R⁸)₂,
- (D)NR⁸C(O)R⁸,

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- 5 (D)NR⁸C(O)OR⁸,
(D)NR⁸C(O)N(R⁸)₂,
(D)NR⁸SO₂R⁸,
(D)SR⁸,
(D)SOR⁸,
10 (D)SO₂R⁸, or
(D)SO₂N(R⁸)₂;

R¹ is independently:

- hydrogen, CONH(C₁-C₈ alkyl), C₁-C₈ alkyl, (D)phenyl, (D)C₃-C₇ cycloalkyl or
15 oxo, provided that oxo is not attached to the same carbon that attached to nitrogen
which forms an amide bond when G is N;

R³ is independently: aryl or thienyl;

- wherein aryl and thienyl are optionally substituted with one to three substituents selected
20 from the group consisting of:
cyano, halo, C₁-C₈ alkyl, (D)C₃-C₇ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkyl
and C₁-C₄ haloalkyloxy;

R⁴ is independently:

- 25 hydrogen, C₁-C₈ alkyl, C(O)R⁸, C(O)OR⁸, C₃-C₇ cycloalkyl or (CH₂)_nO(C₁-C₈
alkyl), wherein n is 2-8;

each R⁸ is independently:

- hydrogen,
30 aryl, wherein aryl being phenyl or naphthyl,
C₁-C₈ alkyl,
C₁-C₄ alkoxy, or

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5 $(\text{CH}_2)_n\text{C}_1\text{-C}_4$ haloalkyl, wherein n is 2-8;

each R^{10} is independently:

hydrogen, $(\text{C}_1\text{-C}_8)$ alkyl, $\text{C}(\text{O})\text{C}_1\text{-C}_8$ alkyl, aryl, or $\text{C}_3\text{-C}_7$ cycloalkyl;

10 each R^{11} is independently:

hydrogen,

$\text{C}_1\text{-C}_8$ alkyl,

aryl,

heteroaryl

15 $(\text{CH}_2)_n\text{N}(\text{R}^8)_2$,

$(\text{CH}_2)_n\text{NR}^8\text{C}(\text{O})\text{C}_1\text{-C}_4$ alkyl,

$(\text{CH}_2)_n\text{NR}^8\text{SO}_2\text{C}_1\text{-C}_4$ alkyl,

$(\text{CH}_2)_n\text{SO}_2\text{N}(\text{R}^8)_2$,

$(\text{CH}_2)_n[\text{O}]_q\text{C}_1\text{-C}_8$ alkyl,

20 $(\text{CH}_2)_n[\text{O}]_q(\text{CH}_2)_n\text{NR}^8\text{COR}^8$,

$(\text{CH}_2)_n[\text{O}]_q(\text{CH}_2)_n\text{NR}^8\text{SO}_2\text{R}^8$,

$(\text{CH}_2)_n[\text{O}]_q$ -heterocyclyl or

$(\text{CH}_2)_n[\text{O}]_q(\text{C}_1\text{-C}_8$ alkyl)-heterocyclyl; and

wherein n is 2-8;

25

each R^{12} is independently:

hydrogen,

$\text{C}_1\text{-C}_8$ alkyl,

$(\text{D})\text{C}_3\text{-C}_7$ cycloalkyl,

30 (D) phenyl

$\text{C}(\text{O})\text{C}_1\text{-C}_8$ alkyl,

$\text{C}(\text{O})$ phenyl,

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- 5 $\text{SO}_2\text{C}_1\text{-C}_8$ alkyl or
 SO_2 -phenyl;

D is a bond or $-(\text{CH}_2)_n$;

n is 0-8;

- 10 p is 0-4; and

 q is 0-1.

2. The compound of Claim 1, wherein the 5- or 6- membered carbocyclyl is cyclohexyl, cyclopentyl, cyclohexenyl, cyclopentenyl or phenyl.

3. The compound of Claim 2, wherein R is independently at each
 15 occurrence: hydrogen, halo

hydroxy, $\text{C}_1\text{-C}_8$ alkyl, $\text{C}_1\text{-C}_8$ alkoxy, $\text{C}_3\text{-C}_7$ cycloalkyl, (D)heterocyclyl, (D) $\text{C}(\text{O})\text{R}^8$,
 (D) $\text{OC}(\text{O})\text{R}^8$, (D) $\text{C}(\text{O})\text{aryl}$, (D) $\text{C}(\text{O})(\text{CH}_2)_n\text{N}(\text{R}^8)_2$, (D) $\text{OC}(\text{O})(\text{CH}_2)_n\text{N}(\text{R}^8)_2$,
 (D) $\text{C}(\text{O})\text{C}_1\text{-C}_4$ alkoxy, (D) $\text{N}(\text{R}^8)_2$, (D) NR^8COR^8 , (D) $\text{NR}^8\text{C}(\text{O})\text{C}_1\text{-C}_4$ alkoxy, (D) SR^8 ,
 (D) SO_2R^8 , (D) $\text{SO}_2\text{N}(\text{R}^8)_2$ or (D) $\text{NR}^8\text{SO}_2\text{R}^8$ where R^8 independently at each

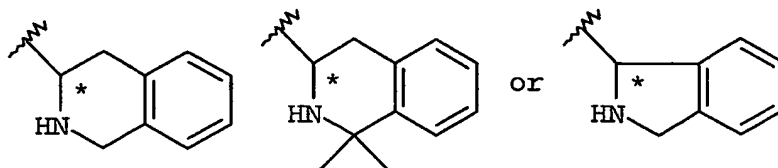
- 20 occurrence being hydrogen, $\text{C}_1\text{-C}_8$ alkyl, $\text{C}_1\text{-C}_8$ alkoxy or phenyl.

4. The compound of Claim 3, wherein R^3 is phenyl optionally para-substituted with chloro, fluoro, bromo, iodo, methoxy, benzyloxy or methyl.

5. The compound of Claim 4, wherein R^3 is phenyl para-substituted with chloro, fluoro or methoxy.

- 25 6. The compound of Claim 5, wherein R^4 is hydrogen.

7. The compound of Claim 6, wherein $-(\text{CH}_2)_n\text{-T}$ is



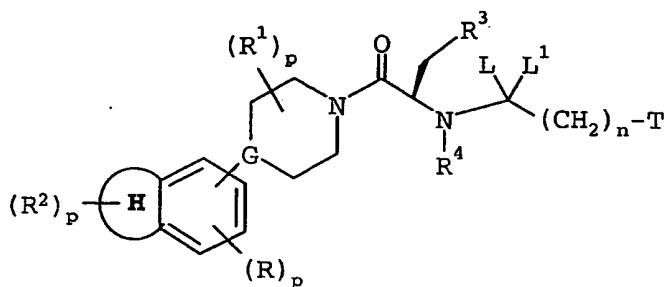
where * denotes a

chiral carbon atom which has a R or S configuration.

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5 8. The compound of Claim 7, wherein L and L¹ are together oxo and the chiral carbon has R configuration.

9. A compound of formula II:



(II)

or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

G is CR¹ or N;

15 L and L¹ are independently hydrogen or together oxo;



is a 5- or 6-membered heterocyclyl,

wherein the heterocyclyl being saturated, partially saturated or aromatic ring, provided that

when G is carbon, the 5-membered nitrogen containing heterocyclyl does not contain a

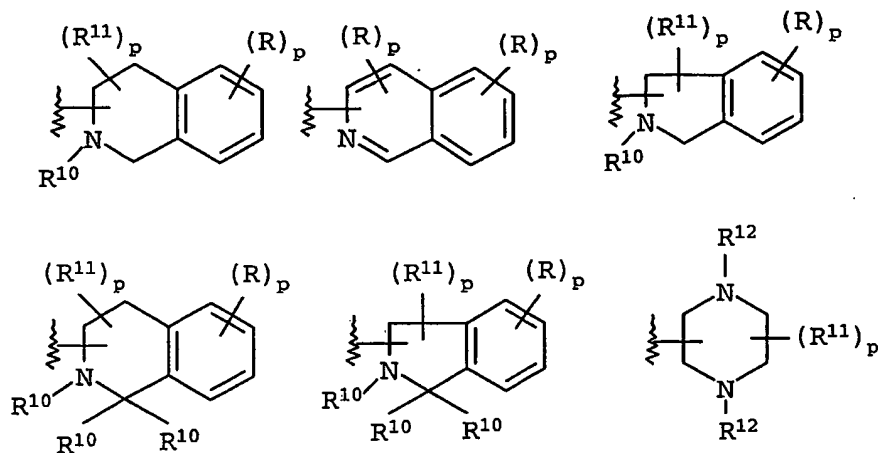
20 carbon-carbon double bond in the ring; and

wherein the heterocyclyl being optionally substituted with one to three substituents

independently selected from R²;

T is:

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5

R is independently:

- hydrogen,
- hydroxy,
- (D)cyano,
- halo,
- C₁-C₈ alkyl,
- C₁-C₈ alkoxy,
- C₃-C₇ cycloalkyl,
- C₁-C₄ haloalkyl,
- (D)heterocyclyl,
- (D)C(O)R⁸,
- (D)C(O)(CH₂)_nN(R⁸)₂,
- C₁-C₈ alkyl-N(R⁸)₂,
- (D)OR⁸,
- (D)OCOR⁸,
- (D)OC(O)N(R⁸)₂,
- (D)N(R⁸)₂,
- (D)NR⁸C(O)R⁸,
- (D)NR⁸C(O)OR⁸,

20

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- 5 (D)NR⁸C(O)N(R⁸)₂,
 (D)NR⁸SO₂R⁸,
 (D)SR⁸,
 (D)SOR⁸,
 (D)SO₂R⁸, or
 10 (D)SO₂N(R⁸)₂;

R¹ is independently:

- hydrogen, CONH(C₁-C₈ alkyl), C₁-C₈ alkyl, (D)phenyl, (D)C₃-C₇ cycloalkyl or
 oxo, provided that oxo is not attached to the same carbon that attached to nitrogen
 15 which forms an amide bond when G is N;

R² is independently:

- hydrogen,
 C₁-C₈ alkyl,
 20 C₃-C₇ cycloalkyl,
 C(O)NH₂,
 C(O)O(C₁-C₆ alkyl),
 C(O)(CH₂)_nN(C₁-C₆ alkyl)₂,
 C(O)-phenyl,
 25 SO₂(C₁-C₆ alkyl),
 SO₂N(C₁-C₆ alkyl)₂,
 SO₂-phenyl or
 C(O)(C₁-C₆ alkyl);

30 R³ is independently: aryl or thienyl;

wherein aryl and thienyl are optionally substituted with one to three substituents selected from the group consisting of:

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- 5 cyano, halo, C₁-C₈ alkyl, (D)C₃-C₇ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkyl
and C₁-C₄ haloalkyloxy;

R⁴ is independently:

- hydrogen, C₁-C₈ alkyl, C(O)R⁸, C(O)OR⁸, C₃-C₇ cycloalkyl or (CH₂)_nO(C₁-C₈
10 alkyl), wherein n is 2-8;

each R⁸ is independently:

- hydrogen,
aryl, wherein aryl being phenyl or naphthyl,
15 C₁-C₈ alkyl,
C₁-C₄ alkoxy, or
(CH₂)_nC₁-C₄ haloalkyl, wherein n is 2-8;

each R¹⁰ is independently:

- 20 hydrogen, (C₁-C₈)alkyl, C(O)C₁-C₈ alkyl, aryl, or C₃-C₇ cycloalkyl;

each R¹¹ is independently:

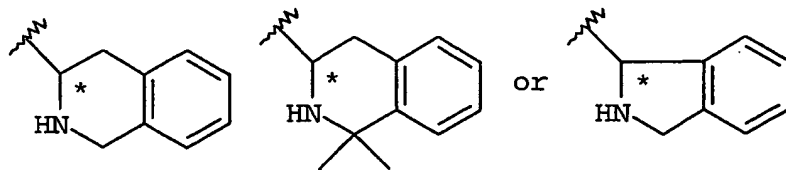
- hydrogen,
C₁-C₈ alkyl,
25 aryl,
heteroaryl
(CH₂)_nN(R⁸)₂,
(CH₂)_nNR⁸C(O)C₁-C₄ alkyl,
(CH₂)_nNR⁸SO₂C₁-C₄ alkyl,
30 (CH₂)_nSO₂N(R⁸)₂,
(CH₂)_n[O]_qC₁-C₈ alkyl,
(CH₂)_n[O]_q(CH₂)_nNR⁸COR⁸,

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- 5 $(\text{CH}_2)_n[\text{O}]_q(\text{CH}_2)_n\text{NR}^8\text{SO}_2\text{R}^8$,
 $(\text{CH}_2)_n[\text{O}]_q$ -heterocyclyl or
 $(\text{CH}_2)_n[\text{O}]_q(\text{C}_1\text{-C}_8 \text{ alkyl})$ -heterocyclyl; and
 wherein n is 2-8;
- 10 each R^{12} is independently:
 hydrogen,
 $\text{C}_1\text{-C}_8$ alkyl,
 (D) $\text{C}_3\text{-C}_7$ cycloalkyl,
 (D)phenyl
15 $\text{C}(\text{O})\text{C}_1\text{-C}_8$ alkyl,
 $\text{C}(\text{O})$ phenyl,
 $\text{SO}_2\text{C}_1\text{-C}_8$ alkyl or
 SO_2 -phenyl;
- 20 D is a bond or $-(\text{CH}_2)_n$;
 n is 0-8;
 p is 0-4; and
 q is 0-1.
- 25 10. The compound of Claim 9, wherein the 5- or 6-membered
 heterocycle contains at least one nitrogen.
11. The compound of Claim 10, wherein the heterocyclyl is
 piperidinyl, pyrrolidinyl, pyrrolinyl, isoxazolyl, oxazolyl, thiazoyl, triazolyl, tetrazolyl,
 thiadiazolyl or oxadiazolyl.
- 30 12. The compound of Claim 11, wherein R^3 is phenyl optionally para-
 substituted with chloro, bromo, benzyloxy, methoxy, fluoro, iodo or methyl.
13. The compound of Claim 12, wherein R^3 is phenyl para-substituted
 with chloro, fluoro or methoxy.
14. The compound of Claim 13, wherein R^4 is hydrogen.

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15. The compound of Claim 14, wherein $-(CH_2)_n-T$ is

where * denotes a

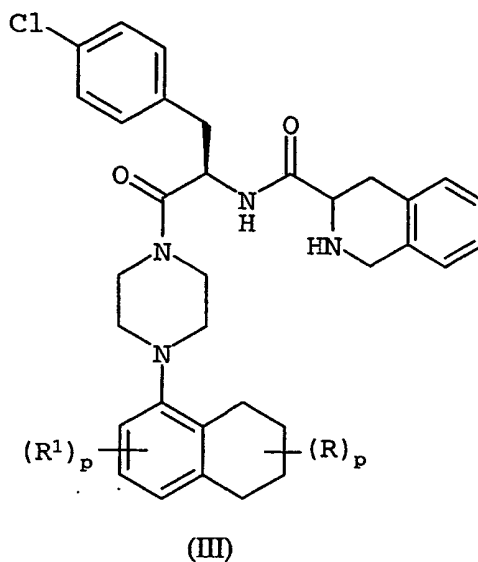
chiral carbon atom which has a R or S configuration.

16. The compound of Claim 15, wherein L and L¹ are together oxo and the chiral carbon has R configuration.

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- 5 17. A compound of formula III,

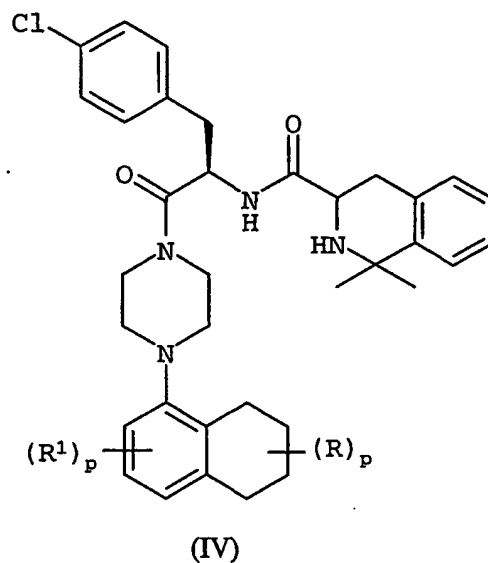


or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

- 10 R is selected from the group consisting of:
 C₁-C₆ alkoxy, NH₂, NH(C₁-C₆ alkyl), N(C₁-C₆ alkyl)₂, NHC(O)O(C₁-C₆
 alkyl), N(CH₃)SO₂(C₁-C₆ alkyl), NHSO₂(C₁-C₆ alkyl), NHC(O)(C₁-C₆ alkyl),
 OC(O)(C₁-C₆ alkyl), OC(O)NH₂, heterocyclyl, C₁-C₆ alkoxy, hydrogen and C₃-
 C₇ cycloalkyl;
- 15 R¹ is selected from the group consisting of:
 halo, C₁-C₆ alkyl, N(C₁-C₆ alkyl)₂, C₁-C₄ haloalkyl, C₁-C₆ alkoxy and
 hydrogen; and
 p is 0-4.

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- 5 18. A compound of formula IV,



or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

R is selected from the group consisting of:

- 10 C₁-C₆ alkoxy, NH₂, NH(C₁-C₆ alkyl), N(C₁-C₆ alkyl)₂, NHC(O)O(C₁-C₆ alkyl), N(CH₃)SO₂(C₁-C₆ alkyl), NHSO₂(C₁-C₆ alkyl), NHC(O)(C₁-C₆ alkyl), OC(O)(C₁-C₆ alkyl), OC(O)NH₂, heterocyclyl, C₁-C₆ alkoxy, hydrogen and C₃-C₇ cycloalkyl;

R¹ is selected from the group consisting of:

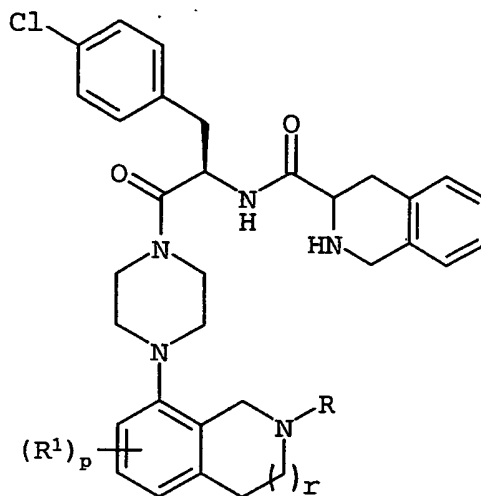
- 15 halo, C₁-C₆ alkyl, N(C₁-C₆ alkyl)₂, C₁-C₄ haloalkyl, C₁-C₆ alkoxy and hydrogen; and

p is 0-4.

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5

19. A compound of formula V,



(V)

or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

R is selected from the group consisting of:

- 10 hydrogen, C₁-C₆ alkyl, C(O)NH₂, C(O)O(C₁-C₆ alkyl), C(O)(CH₂)_nN(C₁-C₆ alkyl)₂, C(O)-phenyl, SO₂(C₁-C₆ alkyl), SO₂N(C₁-C₆ alkyl)₂, SO₂-phenyl, C(O)C₁-C₆ alkyl, C₃-C₇ cycloalkyl and (C₂-C₆ alkyl)N(C₁-C₆ alkyl)₂;

R¹ is selected from the group consisting of:

- 15 halo, C₁-C₆ alkyl, N(C₁-C₆ alkyl)₂, C₁-C₄ haloalkyl, hydrogen and C₁-C₆ alkoxy;

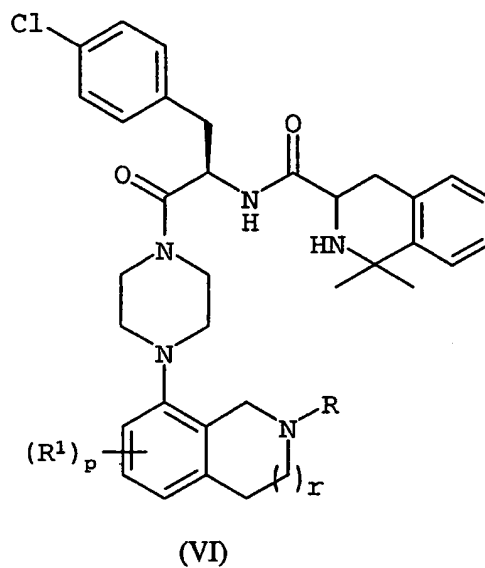
n is 0-8;

p is 0-4; and

r is 0-1.

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- 5 20. A compound of formula VI,



or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

R is selected from the group consisting of:

- 10 hydrogen, C₁-C₆ alkyl, C(O)NH₂, C(O)O(C₁-C₆ alkyl), C(O)(CH₂)_nN(C₁-C₆ alkyl)₂, C(O)-phenyl, SO₂(C₁-C₆ alkyl), SO₂N(C₁-C₆ alkyl)₂, SO₂-phenyl, C(O)C₁-C₆ alkyl, C₃-C₇ cycloalkyl and (C₂-C₆ alkyl)N(C₁-C₆ alkyl)₂;

R¹ is selected from the group consisting of:

- 15 halo, C₁-C₆ alkyl, N(C₁-C₆ alkyl)₂, C₁-C₄ haloalkyl, hydrogen and C₁-C₆ alkoxy;

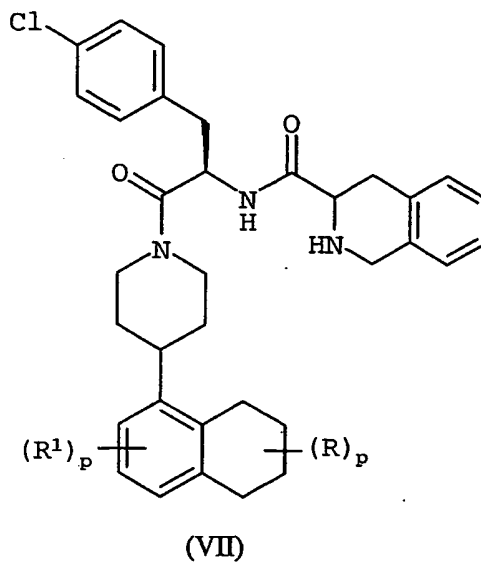
n is 0-8;

p is 0-4; and

r is 0-1.

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- 5 21. A compound of formula VII,



or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

R is selected from the group consisting of:

- 10 C_1 - C_6 alkoxy, NH_2 , $NH(C_1$ - C_6 alkyl), $N(C_1$ - C_6 alkyl) $_2$, $NHC(O)O(C_1$ - C_6 alkyl), $N(CH_3)SO_2(C_1$ - C_6 alkyl), $NHSO_2(C_1$ - C_6 alkyl), $NHC(O)(C_1$ - C_6 alkyl), $OC(O)(C_1$ - C_6 alkyl), $OC(O)NH_2$, heterocyclyl, C_1 - C_6 alkoxy, hydrogen and C_3 - C_7 cycloalkyl;

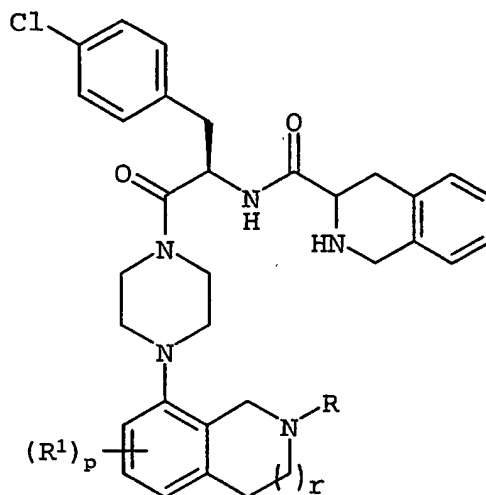
R^1 is selected from the group consisting of:

- 15 halo, C_1 - C_6 alkyl, $N(C_1$ - C_6 alkyl) $_2$, C_1 - C_4 haloalkyl, hydrogen and C_1 - C_6 alkoxy; and

p is 0-4.

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- 5 22. A compound of formula VIII,



(VIII)

or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

R is selected from the group consisting of:

- 10 hydrogen, C₁-C₆ alkyl, C(O)NH₂, C(O)O(C₁-C₆ alkyl), C(O)(CH₂)_nN(C₁-C₆ alkyl)₂, C(O)-phenyl, SO₂(C₁-C₆ alkyl), SO₂N(C₁-C₆ alkyl)₂, SO₂-phenyl, C(O)C₁-C₆ alkyl, C₃-C₇ cycloalkyl and (C₂-C₆ alkyl)N(C₁-C₆ alkyl)₂;

R¹ is selected from the group consisting of:

- 15 halo, C₁-C₆ alkyl, N(C₁-C₆ alkyl)₂, C₁-C₄ haloalkyl, hydrogen and C₁-C₆ alkoxy;

n is 0-8;

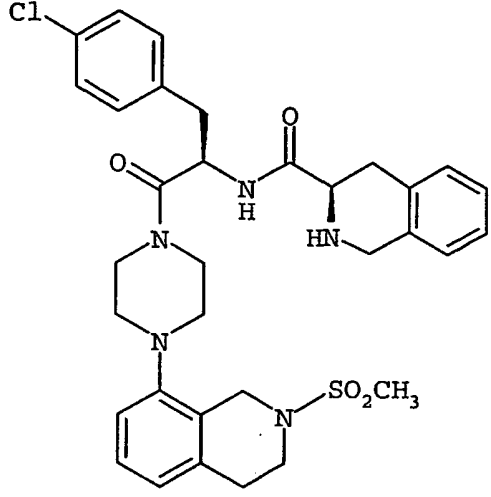
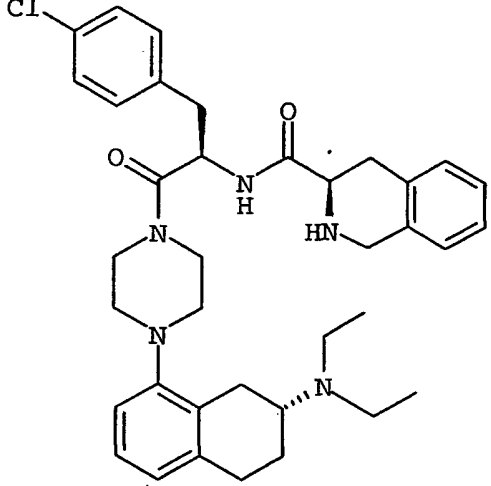
p is 0-4; and

r is 0-1.

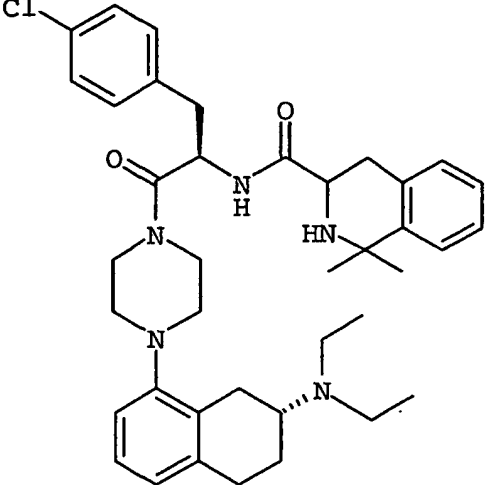
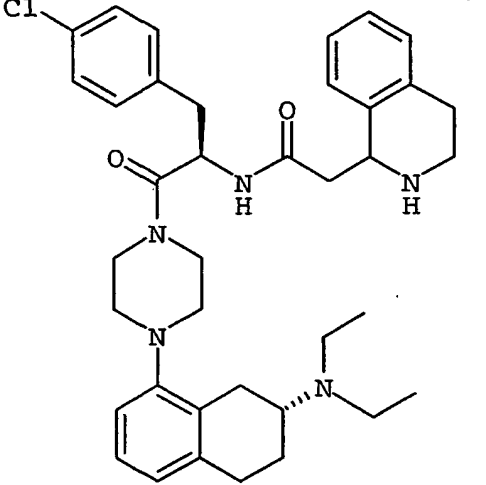
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23. A compound selected from the group consisting of:

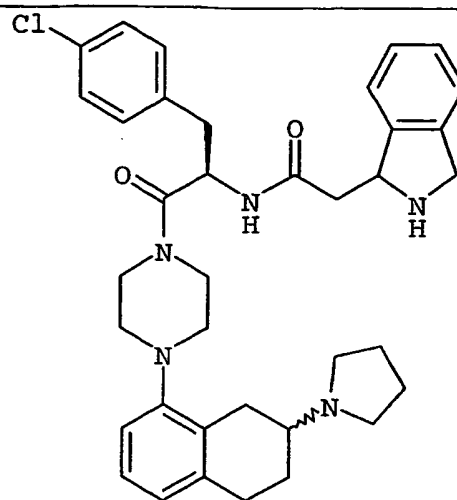
Name of compound	Structure of compound
1,2,3,4-tetrahydro-isoquinoline-3-carboxylic acid {1-(4-chlorobenzyl)-2-[4-(2-methanesulfonyl-1,2,3,4-tetrahydro-isoquinoline-8-yl)-piperazin-1-yl]-2-oxoethyl}-amide	
1,2,3,4-tetrahydro-isoquinoline-3-carboxylic acid {1-(4-chlorobenzyl)-2-[4-(7-diethylamino-5,6,7,8-tetrahydronaphthalen-1-yl)-piperazin-1-yl]-2-oxoethyl}-amide	

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<p>1,1-dimethyl-1,2,3,4-tetrahydro-isoquinoline-3-carboxylic acid {1-(4-chlorobenzyl)-2-[4-(7-diethylamino-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-2-oxo-ethyl}-amide</p>	 <p>and its stereoisomers</p>
<p>N-{1-(4-chloro-benzyl)-2-[4-(7-diethylamino-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-2-oxo-ethyl}-2-(1,2,3,4-tetrahydro-isoquinolin-1-yl)-acetamide</p>	 <p>and its stereoisomers</p>

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N-{1-(4-chloro-benzyl)-2-oxo-2-[4-(7-pyridin-1-yl-5,6,7,8-tetrahydro-naphthalen-1-yl)-piperazin-1-yl]-ethyl}-2-(2,3-dihydro-1H-isoindol-1-yl)-acetamide



and its stereoisomers

5

24. A pharmaceutical composition which comprises a pharmaceutical carrier and at least one compound of formula I or its pharmaceutically acceptable salts or stereoisomers thereof as recited in Claim 1.

25. The pharmaceutical composition of Claim 24, which comprises a second active ingredient selected from the group consisting of: an insulin sensitizer, insulin mimetic, sulfonylurea, alpha-glucosidase inhibitor, HMG-CoA reductase inhibitor, sequestrant cholesterol lowering agent, beta 3 adrenergic receptor agonist, neuropeptide Y antagonist, phosphodiester V inhibitor, and an alpha 2 adrenergic receptor antagonist.

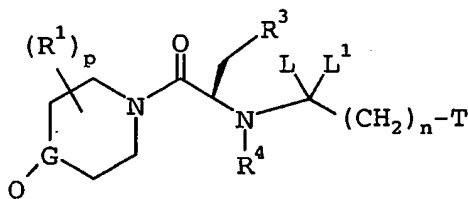
26. A process of making a pharmaceutical composition comprising a compound of formula I or its pharmaceutically acceptable salt or stereoisomers thereof as recited in Claim 1 and a pharmaceutically acceptable carrier.

27. A method of preventing or treating obesity in a mammal comprising the administration of a therapeutically effective amount of the compound of formula I as recited in Claim 1.

28. A method of preventing or treating diabetes mellitus in a mammal comprising the administration of a therapeutically effective amount of the compound of formula I as recited in Claim 1.

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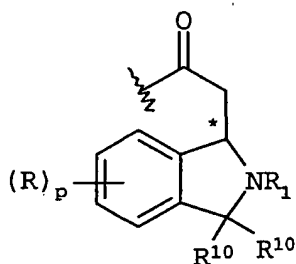
- 5 29. A method of preventing or treating male or female sexual dysfunction in a mammal comprising the administration of a therapeutically effective amount of the compound of formula I as recited in Claim 1.
30. The method of 29, wherein the male or female sexual dysfunction is erectile dysfunction.
- 10 31. A process for preparing a compound of formula I:



(I)

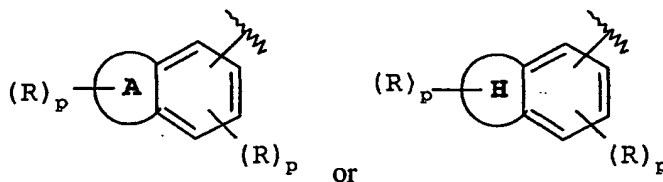
or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

- 15 -CLL¹-(CH₂)ₙ-T is:



wherein R₁ is hydrogen, C₁-C₈ alkyl, Boc, CBZ, Fmoc, phenyl or (C₁-C₈ alkyl)phenyl;

Q represents a moiety:



20

G is CR¹ or N;

-240-

- 5 **A** is a 5- or 6-membered carbocyclyl wherein the carbocyclyl being saturated, partially saturated or aromatic ring, and the carbocyclyl being optionally substituted with one to three substituents independently selected from R;

- H** is a 5- or 6-membered heterocyclyl,
10 wherein the heterocyclyl being saturated, partially saturated or aromatic ring, provided that when G is carbon, the 5-membered nitrogen containing heterocyclyl does not contain a carbon-carbon double bond in the ring; and
wherein the heterocyclyl being optionally substituted with one to three substituents independently selected from R²;

15

R is independently:

- hydrogen,
hydroxy,
(D)cyano,
20 halo,
C₁-C₈ alkyl,
C₁-C₈ alkoxy,
C₃-C₇ cycloalkyl,
C₁-C₄ haloalkyl,
25 (D)heterocyclyl
(D)C(O)R⁸,
(D)C(O)(CH₂)_nN(R⁸)₂,
C₁-C₈ alkyl-N(R⁸)₂,
(D)OR⁸,
30 (D)OCOR⁸,
(D)OC(O)N(R⁸)₂,

-241-

- 5 (D)N(R⁸)₂,
 (D)NR⁸C(O)R⁸,
 (D)NR⁸C(O)OR⁸,
 (D)NR⁸C(O)N(R⁸)₂,
 (D)NR⁸SO₂R⁸,
- 10 (D)SR⁸,
 (D)SOR⁸,
 (D)SO₂R⁸, or
 (D)SO₂N(R⁸)₂;
- 15 R¹ is independently:
 hydrogen, CONH(C₁-C₈ alkyl), C₁-C₈ alkyl, (D)phenyl, (D)C₃-C₇ cycloalkyl or
 oxo, provided that oxo is not attached to the same carbon that attached to nitrogen
 which forms an amide bond when G is N;
- 20 R² is independently:
 hydrogen,
 C₁-C₈ alkyl,
 C₃-C₇ cycloalkyl,
 C(O)NH₂,
- 25 C(O)O(C₁-C₆ alkyl),
 C(O)(CH₂)_nN(C₁-C₆ alkyl)₂,
 C(O)-phenyl,
 SO₂(C₁-C₆ alkyl),
 SO₂N(C₁-C₆ alkyl)₂,
- 30 SO₂-phenyl or
 C(O)(C₁-C₆ alkyl);

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5

R^3 is independently: aryl or thienyl;

wherein aryl and thienyl are optionally substituted with one to three substituents selected from the group consisting of:

10 cyano, halo, C_1 - C_8 alkyl, (D) C_3 - C_7 cycloalkyl, C_1 - C_4 alkoxy, C_1 - C_4 haloalkyl and C_1 - C_4 haloalkyloxy;

R^4 is independently:

hydrogen, C_1 - C_8 alkyl, $C(O)R^8$, $C(O)OR^8$, C_3 - C_7 cycloalkyl or $(CH_2)_nO(C_1$ - C_8 alkyl), wherein n is 2-8;

15

each R^8 is independently:

hydrogen,
aryl, wherein aryl being phenyl or naphthyl,
 C_1 - C_8 alkyl,
20 C_1 - C_4 alkoxy, or
 $(CH_2)_nC_1$ - C_4 haloalkyl, wherein n is 2-8;

each R^{10} is independently:

hydrogen, $(C_1$ - $C_8)$ alkyl, $C(O)C_1$ - C_8 alkyl, aryl, or C_3 - C_7 cycloalkyl;

25

D is a bond or $-(CH_2)_n-$;

n is 0-8;

p is 0-4; and

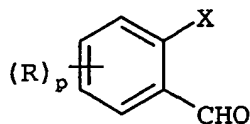
q is 0-1;

30

comprising the steps of:

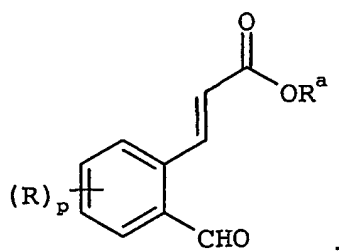
a) reacting a compound having a structural formula 1,

-243-



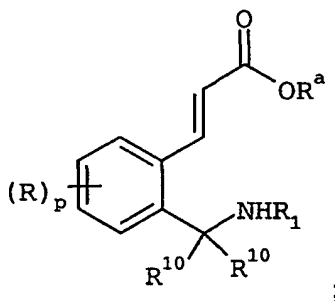
(1)

with $\text{CH}_2\text{CH}=\text{C}(\text{O})\text{OR}^a$ wherein R^a is hydrogen or $\text{C}_1\text{-C}_8$ alkyl and X is halo, in the presence of a catalyst and a base in a suitable organic solvent to give the compound of formula 2,



(2)

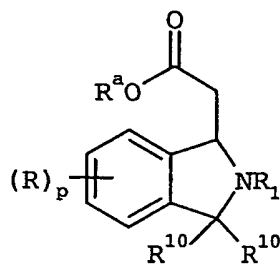
b) reductively aminating the compound of formula 2 in the presence of amine in an acidic condition to give a compound of formula 3,



(3)

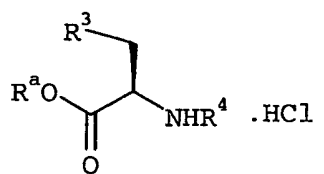
c) cyclizing the compound of formula 3 by Michael addition to give a compound of formula 4 or stereoisomers thereof,

-244-



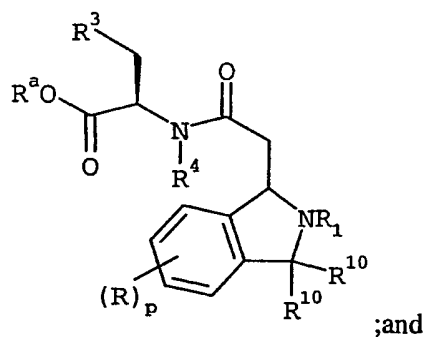
(4)

d) coupling the compound of formula 4 or stereoisomers thereof, wherein R^a of compound 4 is H, with a compound of formula 5,



(5)

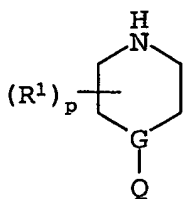
wherein R^a of compound 5 is C₁-C₈ alkyl, to give a compound of formula 6;



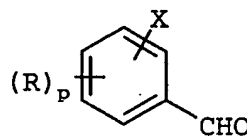
(6)

e) coupling the compound of formula 6, wherein R^a is H, with a compound having a structural,

-245-



5 to afford the compound of formula 1.



32. The process of Claim 31, wherein
is 2-bromobenzaldehydes.

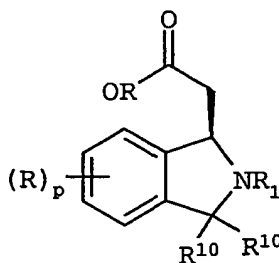
33. The process of Claim 32, wherein $\text{CH}_2\text{CH}=\text{C}(\text{O})\text{OR}$ in Step (a) is
methylacrylate.

10 34. The process of Claim 33, wherein the catalyst in Step (a) is
selected from the group consisting of: $\text{Pd}(\text{Ph}_3\text{P})_2\text{Cl}_2$, $\text{Pd}(\text{Ph}_3\text{P})_4\text{Cl}_2$, $\text{Pd}(\text{Ph}_3\text{P})_4$,
 $\text{Pd}(\text{Ph}_3\text{P})_2\text{Cl}_2/\text{CuI}$, $\text{Pd}(\text{OAc})_2/\text{Ph}_3\text{P}-\text{Bu}_4\text{NBr}$, $\text{Pd}(\text{Ph}_3\text{P})_4\text{Cl}_2/\text{H}_2$ and $\text{Pd}(\text{OAc})_2/\text{P}(\text{O}-$
 $\text{tol})_3$; and wherein the base in Step (a) is NR_3 wherein R is hydrogen or $\text{C}_1\text{-C}_8$ alkyl.

15 35. The process of Claim 34, wherein the amine in Step (b) is selected
from the group consisting of: benzylamine, alpha-methylbenzylamine and BocNH_2 .

36. The process of Claim 35, wherein the Step (b) further comprises
reducing of intermediate imine compound in the presence of reducing agent, the reducing
agent being selected from the group consisting of: NaCNBH_3 , $\text{Na}(\text{OAc})_3\text{BH}$,
 NaBH_4/H^+ , and a combination of Et_3SiH and TFA in CH_3CN or CH_2Cl_2 .

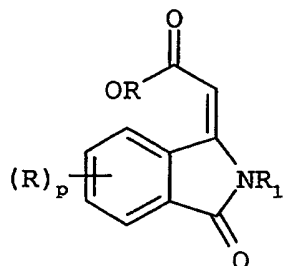
20 37. The process of Claim 31, wherein the stereoisomer of compound of
formula 4 in Step (c) is a compound of formula 4a.



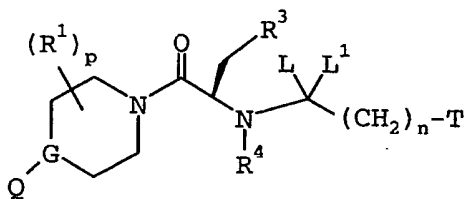
(4a)

-246-

- 5 38. The process of Claim 37, wherein the compound of formula 4a is prepared by asymmetric hydrogenation of a compound having structural formula,



39. The process of Claim 31, wherein the Michael addition in Step (c) is carried out in a basic workup condition.
- 10 40. The process of Claim 31, wherein the Step (e) further comprises deprotecting or protecting of the compound of formula (4) at NR₁.
41. A process for preparing a compound of formula I,

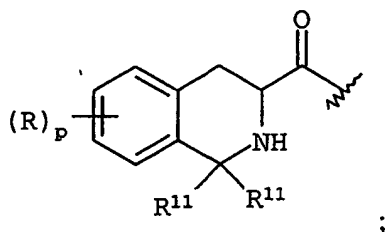


(I)

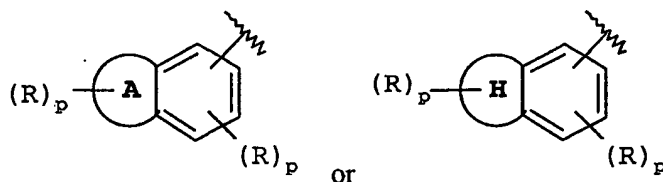
- 15 or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

-247-

5 -CLL¹-(CH₂)_n-T is:



Q represents a moiety:



G is CR¹ or N;

10



is a 5- or 6-membered carbocyclyl wherein the carbocyclyl being saturated, partially saturated or aromatic ring, and the carbocyclyl being optionally substituted with one to three substituents independently selected from R;



15

is a 5- or 6-membered heterocyclyl, wherein the heterocyclyl being saturated, partially saturated or aromatic ring, provided that when G is carbon, the 5-membered nitrogen containing heterocyclyl does not contain a carbon-carbon double bond in the ring; and wherein the heterocyclyl being optionally substituted with one to three substituents

20

independently selected from R²;

R is independently:

hydrogen,

hydroxy,

25

(D)cyano,

halo,

-248-

- 5 C₁-C₈ alkyl,
 C₁-C₈ alkoxy,
 C₃-C₇ cycloalkyl,
 C₁-C₄ haloalkyl,
 (D)heterocyclyl
- 10 (D)C(O)R⁸,
 (D)C(O)(CH₂)_nN(R⁸)₂,
 C₁-C₈ alkyl-N(R⁸)₂,
 (D)OR⁸,
 (D)OCOR⁸,
- 15 (D)OC(O)N(R⁸)₂,
 (D)N(R⁸)₂,
 (D)NR⁸C(O)R⁸,
 (D)NR⁸C(O)OR⁸,
 (D)NR⁸C(O)N(R⁸)₂,
- 20 (D)NR⁸SO₂R⁸,
 (D)SR⁸,
 (D)SOR⁸,
 (D)SO₂R⁸, or
 (D)SO₂N(R⁸)₂;
- 25

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5 R^1 is independently:
hydrogen, $\text{CONH}(\text{C}_1\text{-C}_8 \text{ alkyl})$, $\text{C}_1\text{-C}_8 \text{ alkyl}$, (D)phenyl, (D) $\text{C}_3\text{-C}_7 \text{ cycloalkyl}$ or
oxo, provided that oxo is not attached to the same carbon that attached to nitrogen
which forms an amide bond when G is N;

10 R^2 is independently:
hydrogen,
 $\text{C}_1\text{-C}_8 \text{ alkyl}$,
 $\text{C}_3\text{-C}_7 \text{ cycloalkyl}$,
 $\text{C}(\text{O})\text{NH}_2$,
15 $\text{C}(\text{O})\text{O}(\text{C}_1\text{-C}_6 \text{ alkyl})$,
 $\text{C}(\text{O})(\text{CH}_2)_n\text{N}(\text{C}_1\text{-C}_6 \text{ alkyl})_2$,
 $\text{C}(\text{O})\text{-phenyl}$,
 $\text{SO}_2(\text{C}_1\text{-C}_6 \text{ alkyl})$,
 $\text{SO}_2\text{N}(\text{C}_1\text{-C}_6 \text{ alkyl})_2$,
20 $\text{SO}_2\text{-phenyl}$ or
 $\text{C}(\text{O})(\text{C}_1\text{-C}_6 \text{ alkyl})$;

R^3 is independently: aryl or thienyl;

wherein aryl and thienyl are optionally substituted with one to three substituents selected

25 from the group consisting of:
cyano, halo, $\text{C}_1\text{-C}_8 \text{ alkyl}$, (D) $\text{C}_3\text{-C}_7 \text{ cycloalkyl}$, $\text{C}_1\text{-C}_4 \text{ alkoxy}$, $\text{C}_1\text{-C}_4 \text{ haloalkyl}$
and $\text{C}_1\text{-C}_4 \text{ haloalkyloxy}$;

R^4 is independently:

30 hydrogen, $\text{C}_1\text{-C}_8 \text{ alkyl}$, $\text{C}(\text{O})\text{R}^8$, $\text{C}(\text{O})\text{OR}^8$, $\text{C}_3\text{-C}_7 \text{ cycloalkyl}$ or $(\text{CH}_2)_n\text{O}(\text{C}_1\text{-C}_8 \text{ alkyl})$, wherein n is 2-8;

each R^8 is independently:

hydrogen,

-250-

- 5 aryl, wherein aryl being phenyl or naphthyl,
 C₁-C₈ alkyl,
 C₁-C₄ alkoxy, or
 (CH₂)_nC₁-C₄ haloalkyl, wherein n is 2-8;

- 10 each R¹¹ is independently: hydrogen or (C₁-C₈)alkyl;

D is a bond or -(CH₂)_n-;

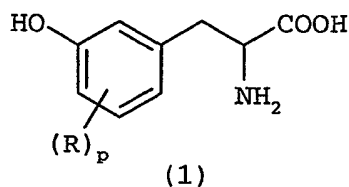
n is 0-8;

p is 0-4; and

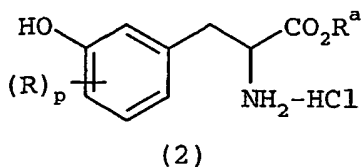
- 15 q is 0-1;

comprising the steps of:

- a) esterifying a compound of formula 1,



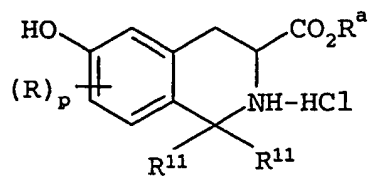
- 20 with an alcohol R^aOH to form a compound of formula 2,



wherein R^a is C₁-C₄ alkyl or (D)phenyl;

- 25 b) reacting a compound of formula 2 with R¹¹COR¹¹ to form a
 compound of formula 3,

-251-

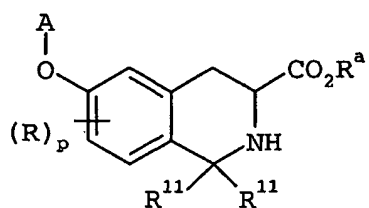


(3)

5

wherein R^{11} is independently hydrogen or C_1 - C_4 alkyl;

c) reacting a compound of formula 3 with an activating group to form a compound of formula 4,

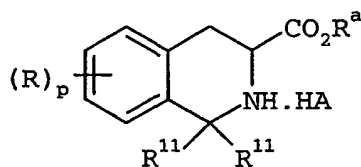


(4)

10

wherein A is an activating group;

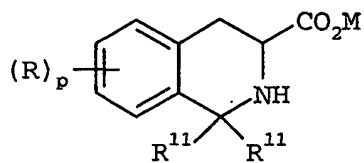
d) deoxygenating the compound of formula 4 by hydrogenation to afford a compound of formula 5,



(5)

15

e) optionally reacting the compound of formula 5 with an inorganic base to form a compound of formula 6,



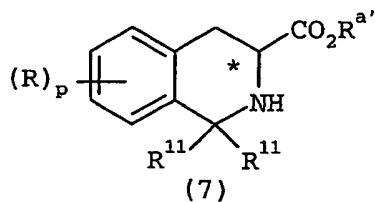
(6)

-252-

5

wherein HA is an acidic and M is a univalent cation;

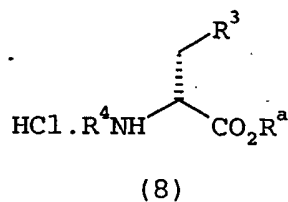
f) resolving the compound of formula 5 or formula 6 to afford a chiral compound of formula 7,



10

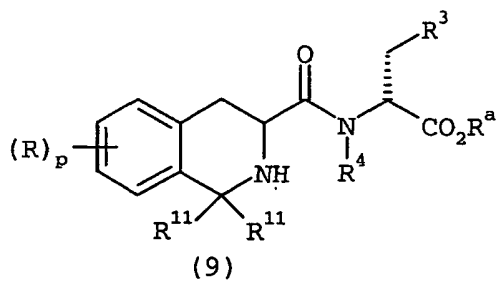
wherein M is hydrogen and R^a is H or R^a;

g) coupling the compound of formula 7 with a compound of formula 8,



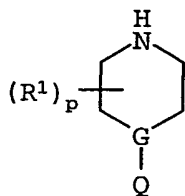
15

to afford a compound of formula 9,



; and

h) coupling the compound of formula 9 with a compound having a

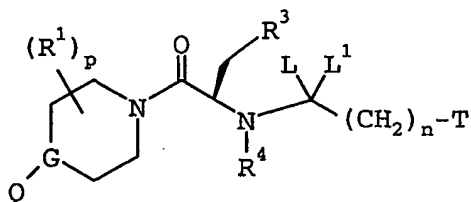


formula,

to afford a compound of formula I.

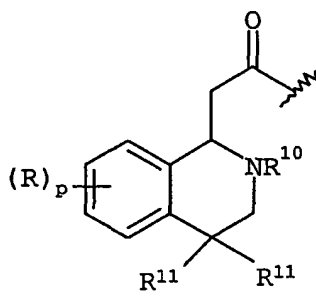
-253-

- 5 42. A process for preparing a compound of formula I:

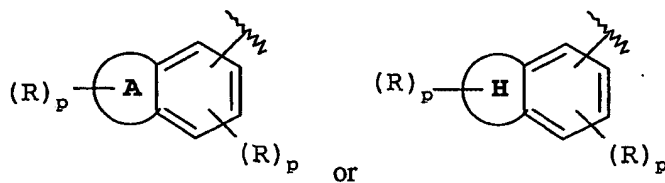


or a pharmaceutically acceptable salts or stereoisomers thereof, wherein

- 10 -CLL¹-(CH₂)ₙ-T is:



Q represents a moiety:



G is CR¹ or N;

- 15 **A** is a 5- or 6-membered carbocyclyl wherein the carbocyclyl being saturated, partially saturated or aromatic ring, and the carbocyclyl being optionally substituted with one to three substituents independently selected from R;

H is a 5- or 6-membered heterocyclyl,

-254-

- 5 wherein the heterocyclyl being saturated, partially saturated or aromatic ring, provided that when G is carbon, the 5-membered nitrogen containing heterocyclyl does not contain a carbon-carbon double bond in the ring; and wherein the heterocyclyl being optionally substituted with one to three substituents independently selected from R²;

10

R is independently:

- hydrogen,
hydroxy,
(D)cyano,
15 halo,
C₁-C₈ alkyl,
C₁-C₈ alkoxy,
C₃-C₇ cycloalkyl,
C₁-C₄ haloalkyl,
20 (D)heterocyclyl
(D)C(O)R⁸,
(D)C(O)(CH₂)_nN(R⁸)₂,
C₁-C₈ alkyl-N(R⁸)₂,
(D)OR⁸,
25 (D)OCOR⁸,
(D)OC(O)N(R⁸)₂,
(D)N(R⁸)₂,
(D)NR⁸C(O)R⁸,
(D)NR⁸C(O)OR⁸,
30 (D)NR⁸C(O)N(R⁸)₂,
(D)NR⁸SO₂R⁸,
(D)SR⁸,

-255-

- 5 (D)SOR⁸,
(D)SO₂R⁸, or
(D)SO₂N(R⁸)₂;

R¹ is independently:

- 10 hydrogen, CONH(C₁-C₈ alkyl), C₁-C₈ alkyl, (D)phenyl, (D)C₃-C₇ cycloalkyl or
oxo, provided that oxo is not attached to the same carbon that attached to nitrogen
which forms an amide bond when G is N;

R² is independently:

- 15 hydrogen,
C₁-C₈ alkyl,
C₃-C₇ cycloalkyl,
C(O)NH₂,
C(O)O(C₁-C₆ alkyl),
20 C(O)(CH₂)_nN(C₁-C₆ alkyl)₂,
C(O)-phenyl,
SO₂(C₁-C₆ alkyl),
SO₂N(C₁-C₆ alkyl)₂,
SO₂-phenyl or
25 C(O)(C₁-C₆ alkyl);

R³ is independently: aryl or thienyl;

wherein aryl and thienyl are optionally substituted with one to three substituents selected
from the group consisting of:

- 30 cyano, halo, C₁-C₈ alkyl, (D)C₃-C₇ cycloalkyl, C₁-C₄ alkoxy, C₁-C₄ haloalkyl
and C₁-C₄ haloalkyloxy;

-256-

5 R^4 is independently:

hydrogen, C_1 - C_8 alkyl, $C(O)R^8$, $C(O)OR^8$, C_3 - C_7 cycloalkyl or $(CH_2)_nO(C_1$ - C_8 alkyl), wherein n is 2-8;

each R^8 is independently:

10 hydrogen,
aryl, wherein aryl being phenyl or naphthyl,
 C_1 - C_8 alkyl,
 C_1 - C_4 alkoxy, or
 $(CH_2)_nC_1$ - C_4 haloalkyl, wherein n is 2-8;

15

each R^{10} is independently:

hydrogen, $(C_1$ - $C_8)$ alkyl, $C(O)C_1$ - C_8 alkyl, aryl, C_3 - C_7 cycloalkyl, or a protecting group selected from Boc, CBZ or FMOC;

20 each R^{11} is independently: hydrogen or $(C_1$ - $C_8)$ alkyl;

D is a bond or $-(CH_2)_n$;

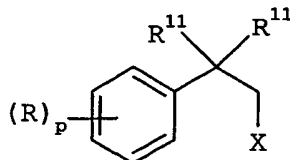
n is 0-8;

p is 0-4; and

25 q is 0-1;

comprising the steps of:

a) reacting a compound formula 1:

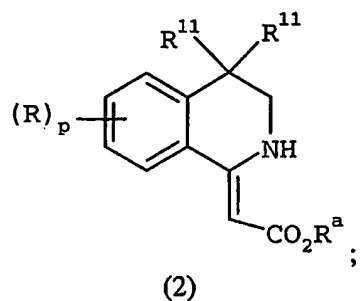


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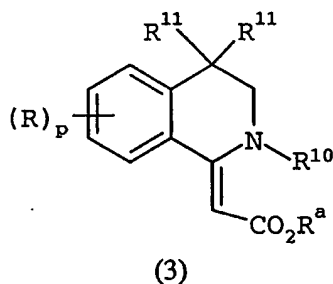
(1)

-257-

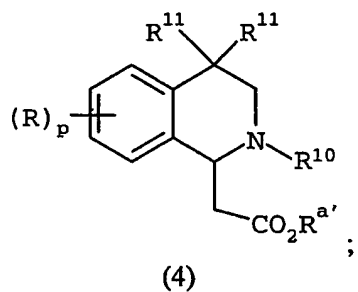
- 5 wherein X is halo, and R^{11} is independently, hydrogen or C1-C4 alkyl, with $CNCH_2CO_2R^a$ wherein R^a is C1-C8 alkyl or benzyl to afford a compound of formula 2:



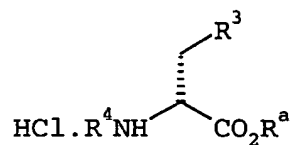
- 10 b) protecting the compound of formula 2 to form the compound of formula 3:



- 15 c) hydrogenating the compound of formula 3 to afford a compound of formula 4:



- 20 d) coupling the compound of formula 4 wherein $R^{a'}$ is hydrogen or R^a , with a compound of formula 5,

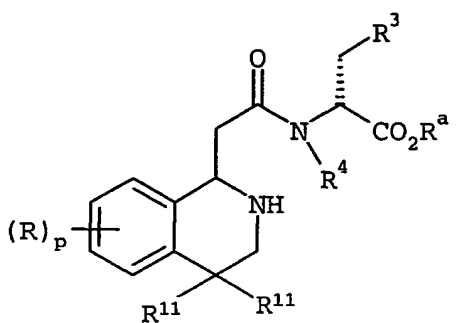


-258-

5

(5)

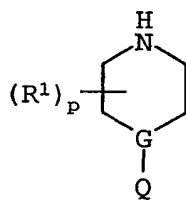
to afford a compound of formula 6,



(6)

10

e) coupling the compound of formula 6 with a compound having a



formula,

to afford a compound of formula I.

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 02/00516

A. CLASSIFICATION OF SUBJECT MATTER

IPC 7 C07D401/12 C07D401/14 A61P15/10 A61K31/44 A61K31/495

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 7 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, WPI Data, PAJ, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P,X	US 6 294 534 B1 (PATCHETT ARTHUR A ET AL) 25 September 2001 (2001-09-25) cited in the application * see claim 1, definition of the spiro part * the whole document	1-42
P,X	WO 01 70337 A (MERCK & CO INC ;NARGUND RAVI P (US); PALUCKI BRENDA L (US)) 27 September 2001 (2001-09-27) cited in the application * see the definition of the spiro part * the whole document	1-42
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☒ Further documents are listed in the continuation of box C.☒ Patent family members are listed in annex.

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Date of the actual completion of the international search

12 June 2002

Date of mailing of the international search report

19/06/2002

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C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 01 70708 A (POLLARD PATRICK G ;LAI YINGJIE (US); YE ZHIXIONG (US); GUO LIANGQI) 27 September 2001 (2001-09-27) cited in the application *see the definition of X and Y * the whole document ----	1-42
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Information on patent family members

International Application No

PCT/US 02/00516

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